

DRESLER, P.I.

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AUTHORS: Bresler, P.I. and Ruzin, B.N.

S/051/50/009/01/004/031
E201/E691

TITLE: The Optico-Acoustic Effect in the Visible and Ultraviolet Regions, and Its Relationship with Photochemical Reactions in Gases

PERIODICAL: Optika i spektroskopiya, 1960, Vol 9, Nr 1, pp 22-25 (USSR)

ABSTRACT: The optico-acoustic effect was discovered in the infrared region. Recently Gerlovin (Ref 1) reported observations of the optico-acoustic effect in the ultraviolet region in nitrogen, oxygen and acetylene. The present paper reports the existence of the optico-acoustic effect in the near ultraviolet region in chlorine and in the visible region in nitrogen dioxide. The optico-acoustic effect was detected by means of apparatus (Fig 1) consisting of a typical optico-acoustic gas analyser and including appropriate light sources. The ultraviolet sources were lamps UFO and FRK-4; the visible source was a 12V, 30W incandescent lamp. The authors obtained also the "gas characteristics" of a prototype optico-acoustic gas analyser

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E201/E691

The Optico-Acoustic Effect in the Visible and Ultraviolet Regions, and Its
Relationship with Photochemical Reactions in Gases

for chlorine. It was found that the optico-acoustic effect was strongly enhanced on addition of hydrogen to chlorine contained in an optico-acoustic chamber. This was due to an additional optico-acoustic effect produced by a photochemical reaction of chlorine and hydrogen. There are 3 figures and 1 Soviet reference.

SUBMITTED: October 29, 1959

W

Card 2/2

S/051/62/013/003/003/012
E032/E514

AUTHOR: Bresler, P.I.

TITLE: An approximate method of determination of the integral intensity and half-width of lines in the vibration-rotation absorption bands of gases

PERIODICAL: Optika i spektroskopiya, v.13, no.3, 1962, 313-316

TEXT: It is pointed out that the relation between the magnitude of the optico-acoustic effect and the properties and density of the gas under investigation which fills the chamber of an optico-acoustic detector of radiation may be used in an approximate determination of the integral intensity and half-width of rotational lines. This method is said to have been first described by M. L. Veyngerov and P. V. Slobodskaya (Izv. AN SSSR, ser. fiz., 11, 420, 1947) and was discussed theoretically by the author in a previous paper (Opt. i spektr., 7, 616, 1959). It is noted that the optico-acoustic detector has a basic advantage as compared with the method described by E. B. Wilson and A.J. Wells (J. Chem. Phys., 14, 578, 1946). The latter method involves the spectral decomposition of the radiation and subsequent graphical

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An approximate method of ...

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integration, while the optico-acoustic method does not require this spectral decomposition. It is now pointed out that there is a range of values for the equivalent thickness of the gas under investigation in the optico-acoustic detector for which the amplitude of the acoustic vibrations is proportional to the square-root of this thickness. By determining the minimum and maximum values of the thickness between which the square-root dependence holds, it is possible to determine the integral intensity and half-width (α and δ respectively). In fact, these two quantities are given by:

$$\alpha \approx \frac{4.96 \cdot B}{\sqrt{m}} \cdot \frac{1}{\sqrt{\frac{w_{\min}}{w_{\max}}}} \quad (5)$$

$$\delta \approx 0.146 \cdot d \cdot \sqrt{\frac{w_{\min}}{w_{\max}}} \quad (6)$$

Card 2/3

An approximate method of ...

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E032/E514

where B is the rotational constant of the molecule and $m = Bhc/kT$. Trial experiments based on this method and designed to determine the integral intensity of the 14.8μ band of CO_2 have shown that the method is only suitable for the determination of rough values of α and δ . This might be useful in studying bands which are not well known and is, therefore, of interest in gas-analytic practice. There are 2 figures.

SUBMITTED: June 21, 1961

Card 3/3

BENSIER, S.D., inzh.; KAVALERCHIK, M.Ya., inzh.

Air purification and recirculation in picking shops. Tekst. prom.
18 no.6:32-33 Je '58, (MIRA 11:7)
(Textile factories--Heating and ventilation)

PROCESSES AND PROPERTIES 6001

The effect of the addition of salts on swelling in willow tanner's liquor. S. Bressler. *Vestnik Kozhevnikov Prom.* Torgor, 1930, 216 III; *Chem. Zentr.* 1932, II, 3809. - The effect of the addition of various salts to willow tanner's liquor on the swelling of gelatin and unhaired hide was investigated for liquors of 1°, 2° and 3°Bé. strength and salt concns. of 0.05, 0.1 and 0.15 M. In the pure 1°Bé. liquor contg. no salt the increase in wt. of the gelatin in 21 hrs. was 330%; in the presence of NaCl, 018%; NaF, 020%; urea, 634%; NaHSO₄, 640% and NaSO₄, 104%. In all cases the swelling decreased with increasing strength of the liquor and increased with salt concn. The following were investigated in liquors of 1°Bé. strength: K₂SO₄, NH₄Cl, NaHPO₄, FeSO₄, CuSO₄, FeNH₄(SO₄)₂, Cr₂(SO₄)₃, K₂CrO₇, K₄Fe(CN)₆ and K₄Fe(CN)₆. A reduction of the swelling nitrates, Co and Zn acetates. A reduction of the swelling was observed in the presence of K₂CrO₇, nitrobenzene and nitrotoluene. The same effect was observed on unhaired hides in the case of K₂CrO₇ and nitrobenzene.

M. G. Moore

ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION

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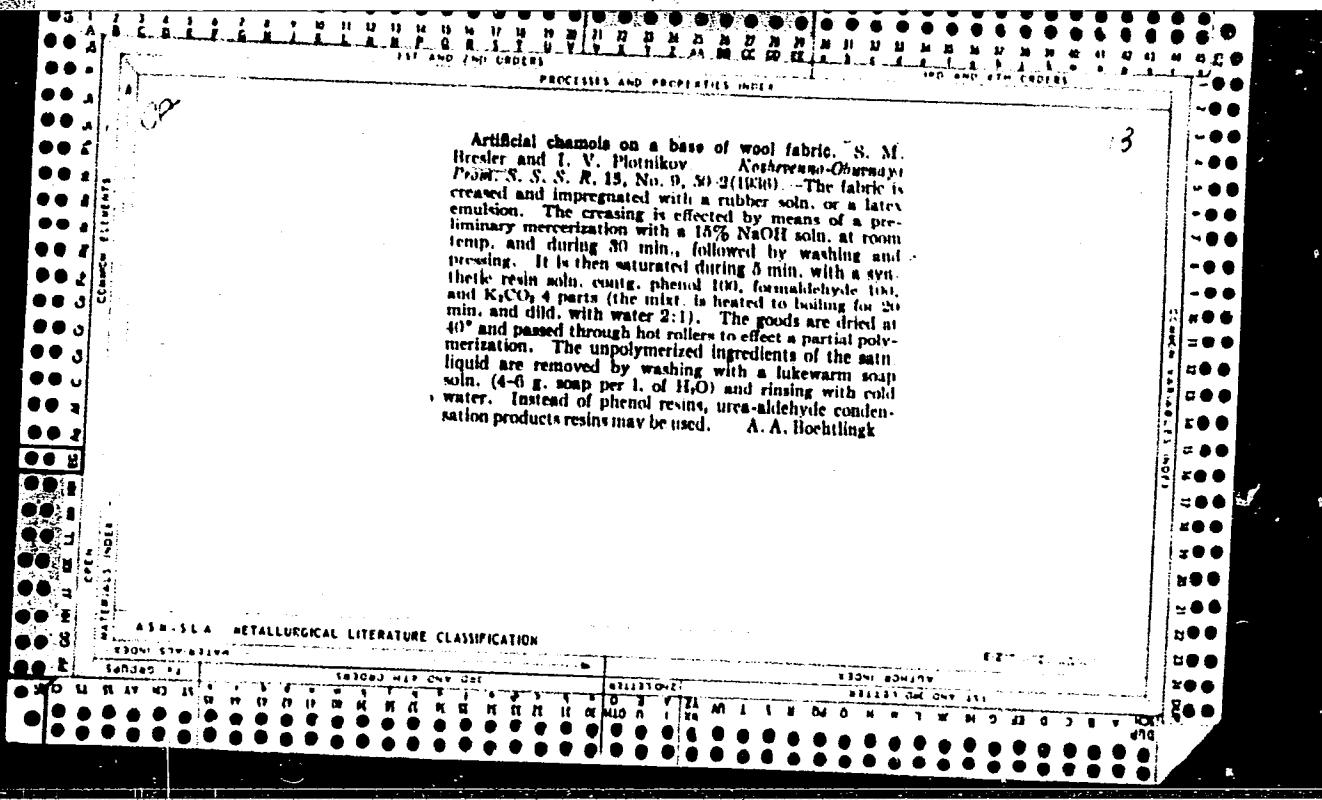
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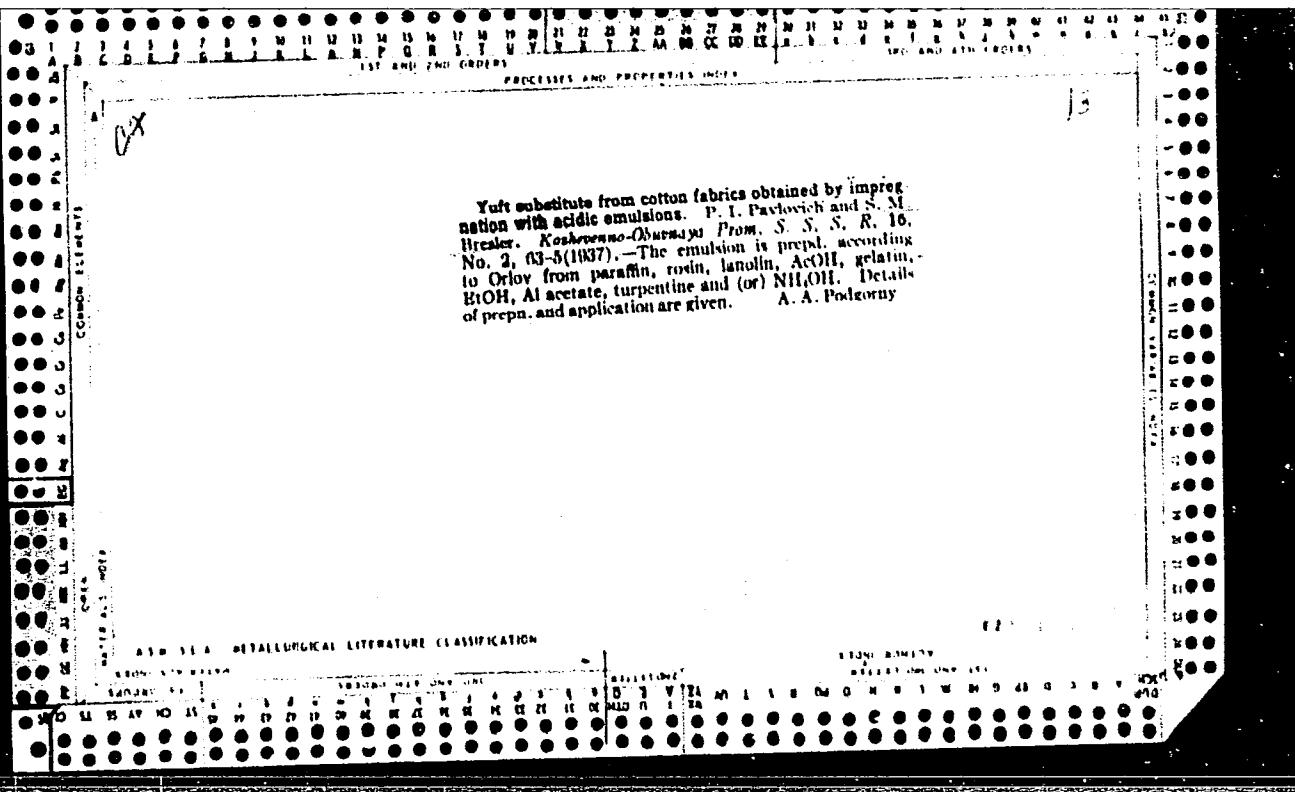
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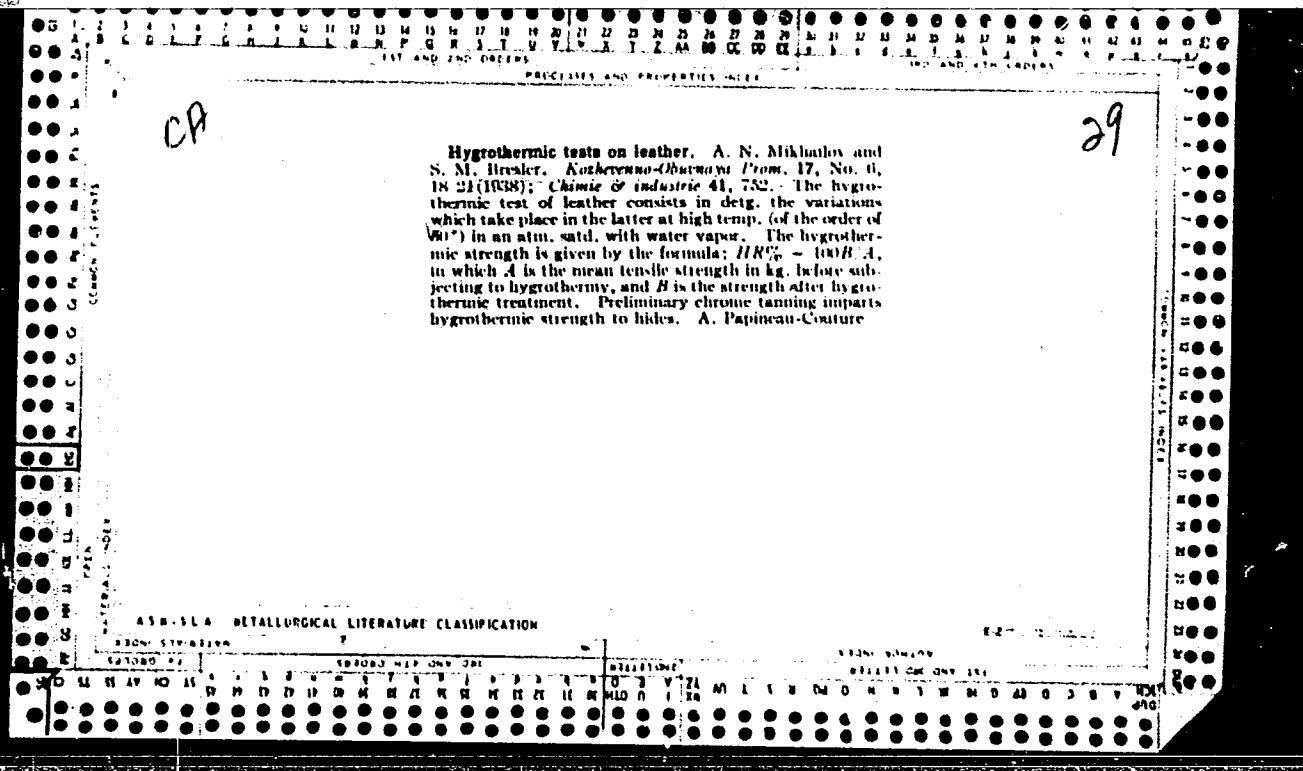
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✓ Use of urotropine in mineral and chrome-vegetable tanning. S. M. Bresler and A. N. Mikhailev. *Zhurnal Pochvovedeniya*, 1960, 14(1), 1-10. Comparison of the interaction between urotropine and Cr and Al salts in the complex formation of protein with gelatin. The interaction of urotropine and Cr or Al salt with gelatin is reversible. The chrome urotropine complex is stable at pH 4-6, further acidic or alkaline causes destruction of the complex. Urotropine acts as a buffer when added to Al-salt soln. Urotropine raises the shrinkage temp. of both chrome and chrome-vegetable tanned leather. P. J. Kanach.

BRESLER, S.

Mikhailov, A. Utilization of urotropine in mineral and chromium tannin. Tr. from the Russian. p. 15.
LEKA PROMISHLENOST, Sofiya, Vol. 4, no. 2, 1955.

SO: Monthly List of East European Accessions, (EEAL), LC, Vol. 4, no. 10, Oct. 1955,
Uncl.

BRESLER, S.M., kandidat tekhnicheskikh nauk

Effect of aluminum salts in chrome tanning. Leg. prom. 15 no.4:
44-45 Ap '55.
(MLRA 8:?)

(Tanning)

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

BRESTER S.M.

BRESTER, S.M., kand.tekhn.nauk

Method of determining the tanning efficiency of chromium salts.
Leg.prom.17 no.9:32-33 S '57. (MIRA 10:12)
(Chromium salts) (Tanning)

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

BRESLER, S.M., kand. tekhn. nauk; MIKHAYLOV, A.N., doktor tekhn. nauk prof.

Acidity conditions in chrome tanning. Kozh.-obuv. prom. no. 8:15-17
Ag '59. (Tanning) (MIRA 13:1)

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

BRESIER, S. M., kand.tekhn.nauk; MIKHAYLOV, A.N., doktor tekhn.nauk, prof.

Tanning chrome pig leather with chrome emulsions. Kozh.-obuv.
prom. 2 no.4:12-15 Ap '60. (MIRA 13:9)
(Tanning)

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

BRESLER, S.M.; MIKHAYLOV, A.N.

Effect of the methods of the basicity regulation of chromium
salt solutions on their tanning properties. Kozh.-obuv.prom.
4 no.6:19-23 Je '62. (MIRA 15:6)
(Tanning materials)

BRESLER, S.M.; MIKHAYLOV, A.N.

Improving the chrome emulsion method of leather tanning. Kozh.-
obuv.prom. 4 no.8:32-34 Ag '62. (MIRA 15:8)
(Tanning)

BRESLER, S.M.; MIKHAYLOV, A.N.

Effect of the tanning temperature on the stability of the
chromium-collagen bond in leather washing with water. Kozh.-
obuv. prom. 5 no.11:29-32 N '63. (MIRA 17:1)

BRESLER, S.M.; MIKHAYLOV, A.N.; ROGATYKH, N.I.

Increased resistance to the washing out of leather fixed
chromium compounds. Nauch.-issl. trudy TSNIKP no. 20:33-36
'63

(MIRA 18:1)

BRESLER, S.M.; MIKHAYLOV, A.N.

Changes in the composition of chromium complexes occurring
during tanning. Nauch.-issl. trudy TSNIKP no.33:24-30 1963

(MIRA 18:1)

PHASE I BOOK EXPLOITATION

SOV/6568

Bresler, Semen Yefimovich

Vvedeniye v molekulyarnuyu biologiyu (Introduction to Molecular Biology)
Moscow, Izd-vo AN SSSR, 1963. 519 p. Errata slip inserted. 7500 copies
printed.

Sponsoring Agency: Akademiya nauk SSSR. Institut vysokomolekulyarnykh
soyedineniy.

Resp. Ed.: S. A. Neyfakh, Doctor of Biological Sciences, Professor; Ed. of
Publishing House: A. A. Frolov; Tech. Ed.: R. A. Zamarayeva.

PURPOSE: This book is intended for scientists and graduate students in chemistry,
physics, and biology.

COVERAGE: The book reviews modern achievements in a comparatively new
field of science, devoted to the study of life on a molecular level.

Card 1/8

Introduction to Molecular Biology

SOV/6568

level. The author defines the role, and the place of molecular biology in natural science and shows the direction of past and present trends in the study of high molecular compounds occurring in nature (proteins and nucleic acids) and the progress being made in the study of their structure, functions, and synthesis in the cell. There are 208 references.

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Card 2/6

BRESLER, S.Ye.; PYRKOV, L.M.; FRENKEL', S.Ya.

Sedimentation of graft copolymers in a density gradient. Approach
to equilibrium, selective solvation, and polydispersity of composition.
Vysokom. soed. 5 no.9:1315-1320 S '63. (MIRA 17:1)

1. Institut vysokomolekularnykh soyedineniy AN SSSR.

BRESLER, S.Ye.; POPOV, A.G.

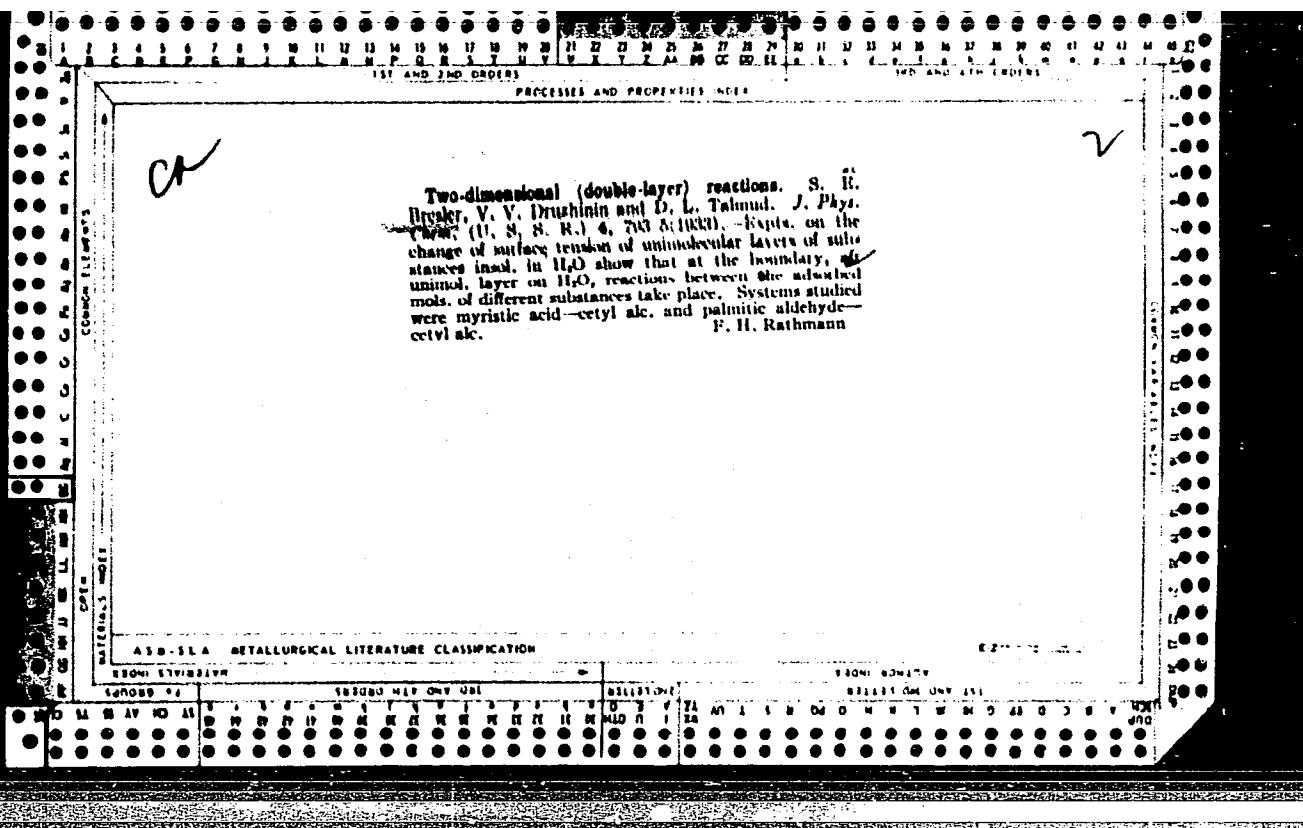
Gas-liquid chromatography. Part 3. Zhur. fiz. khim. 37 no.5:
1178-1182 My '63.
(MIRA 17:1)

1. Institut vysokomolekulyarnykh soyedineniy AN SSSR.

BRESLER, S.Ye.; PERUMOV, D.A.

Mutagenesis on isolated DNA induced by ultraviolet radiation
and chemical agents. Dokl. AN SSSR 158 no.4:967-969 O '64.

(MIRA 17:11)
1. Institut vysokomolekulyarnykh soyedineniy AN SSSR. Pred-
stavлено академиком A.N. Tereninym.



The depth effect of orientating forces and the stability of foams and emulsions. S. E. Bresler and D. L. Talmud. — *J. Phys. Chem.* (U. S. S. R.) 47, 790-801 (1933). — Exptn. with 3 to 25% aq. solns. of KCl, KBr, KI, K₂SO₄, K₄Fe(CN)₆, KCNS and NaCl in C₆H₆ emulsions were made to measure the stability of elementary foams and emulsions formed by non-aq. solns. of non-polar substances and solns. of electrolytes, and are summarized in several tables. The results substantiate the views that quasi-crystal complexes are oriented by the effect of the boundary layer and that in such case the hydration of polar groups must exert an additional depth effect in some sizes of complexes. The surface tensions at 20° for solns. of biphenyl in C₆H₆ were 0.65, 29.20; 20%; 29.43; 30%; 31.39; 30%; 32.12 and for naphthalene in C₆H₆ 0%; 21.17; 30%; 21.81 dynes/cm.
p. H. Rathmann

The mobility of the molecules in the surface layer. S. E. Bensusan, B. A. Talmud and D. L. Talmud. *Physik Z.*, **Symposium 6**, 604-72 (1963).—Layers of acrylic acid and other acids were filtered through a "hot capillary" onto a water surface. The curve of surface tension variation with time shows a period of slow variation (extension against mobility). A relation between rate of flow, viscosity of the film, and the resistance of the base is indicated. Louis Goldman

Louis Goldman

The structure of the nucleus layer of a liquid. V. K. Borkovskii, S. B. Ryazan', I. L. Zelenin and E. A. Strel'tsov. *Physik Z. Sowjetunion* 4, 573-584 (1953).—The orienting effect of the surface extends to within 10^{-3} cm as shown by the thickness of bubbles in parallel 2-3° above its m. p. and the thickness of a paraffin film that prevents the coalescence of 3 Hg layers. Electrostatic diagrams for Hg above a quasi-cryst. structure for the surface.

cryst. structure for
Louis Goldman

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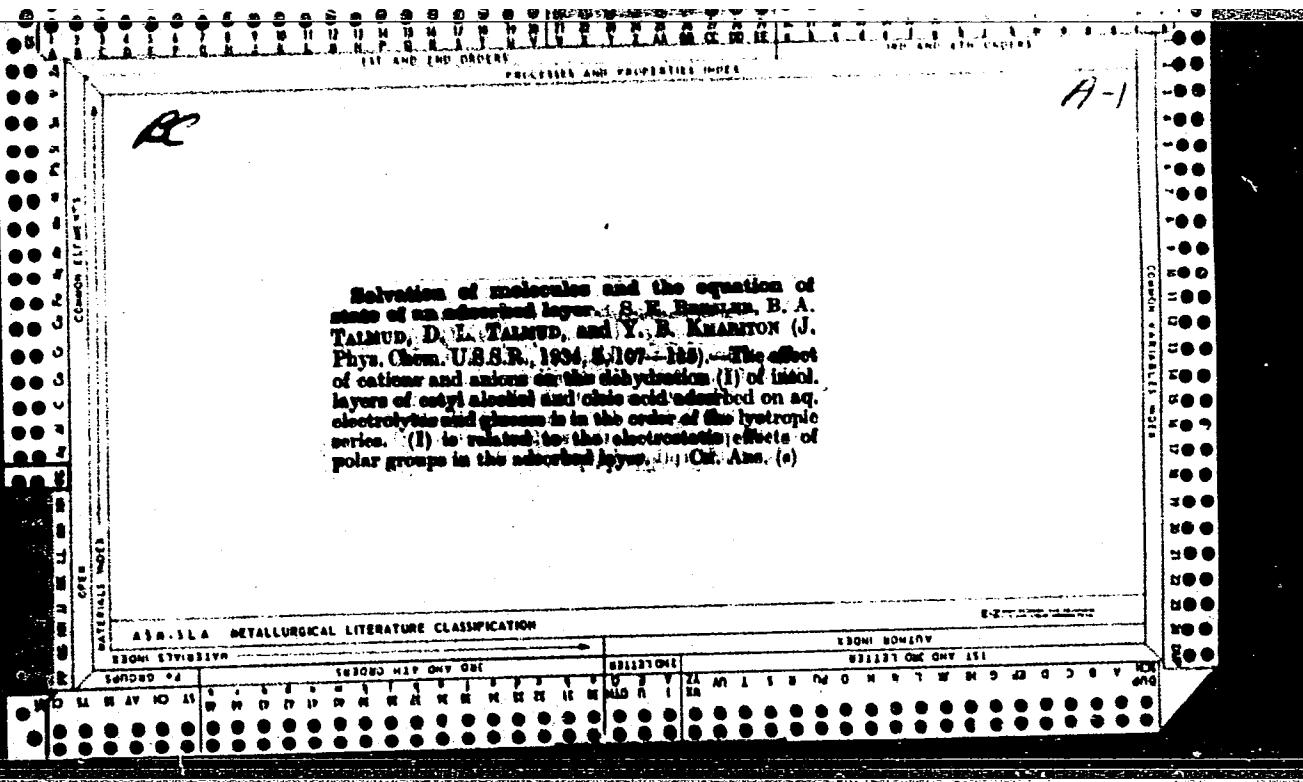
Gelatinized emulsions. S. E. Breiter, M. Kats and L. Kremnev. *Physik. Z. Sowjetunion* 6, 149-153 (1933).
The prepa. of gelatinized emulsions is described. They contain up to 90% by vol. of an org. liquid which in the emulsion has a frothy structure. Louis Goldmark

The recrystallization of dispersed quartz. S. E. Breiter, B. A. Strauf and I. L. Zelmanov. *Physik. Z. Sowjetunion* 6, 150-153 (1933).—A film of amorphous quartz formed by condensation of the vapor recrystallizes at about 400° as compared with 1000° for the recrystall. of quartz glass. Louis Goldmark

Cancer Element

Cancer Agent

ASA-LIA METALLURGICAL LITERATURE CLASSIFICATION

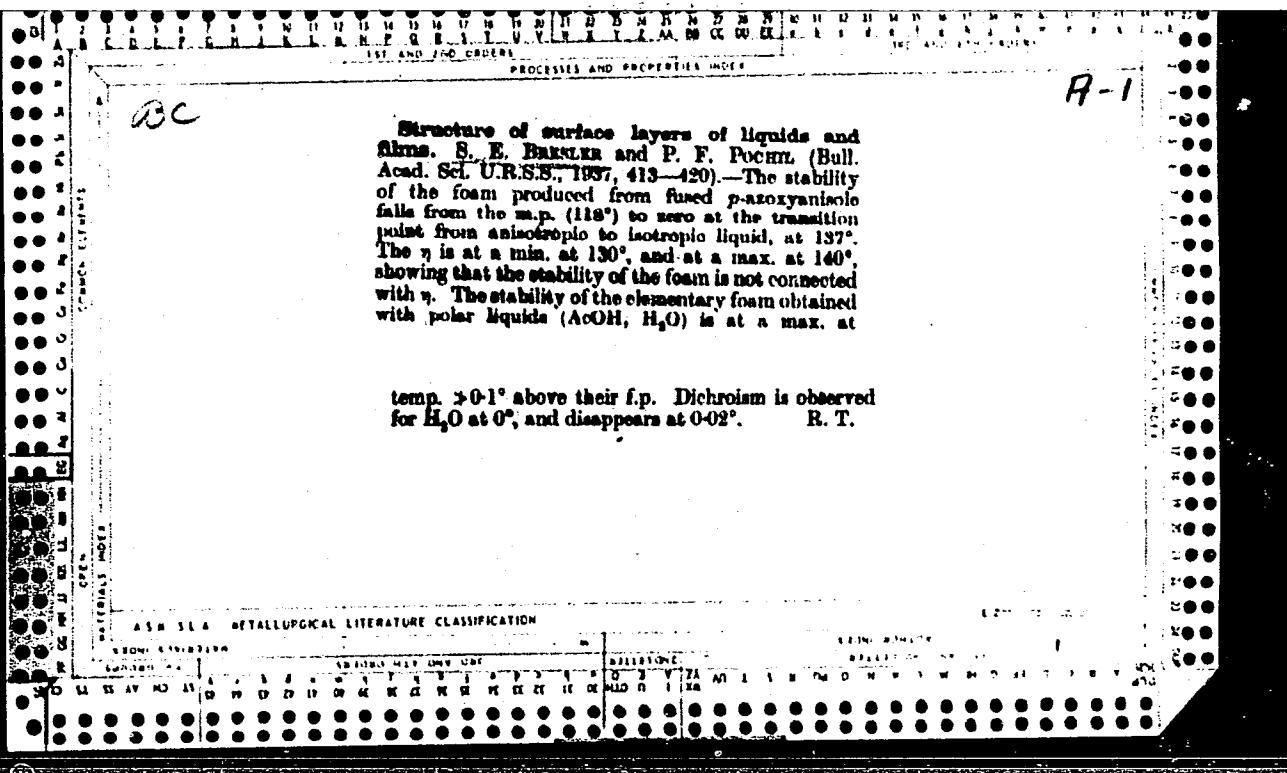


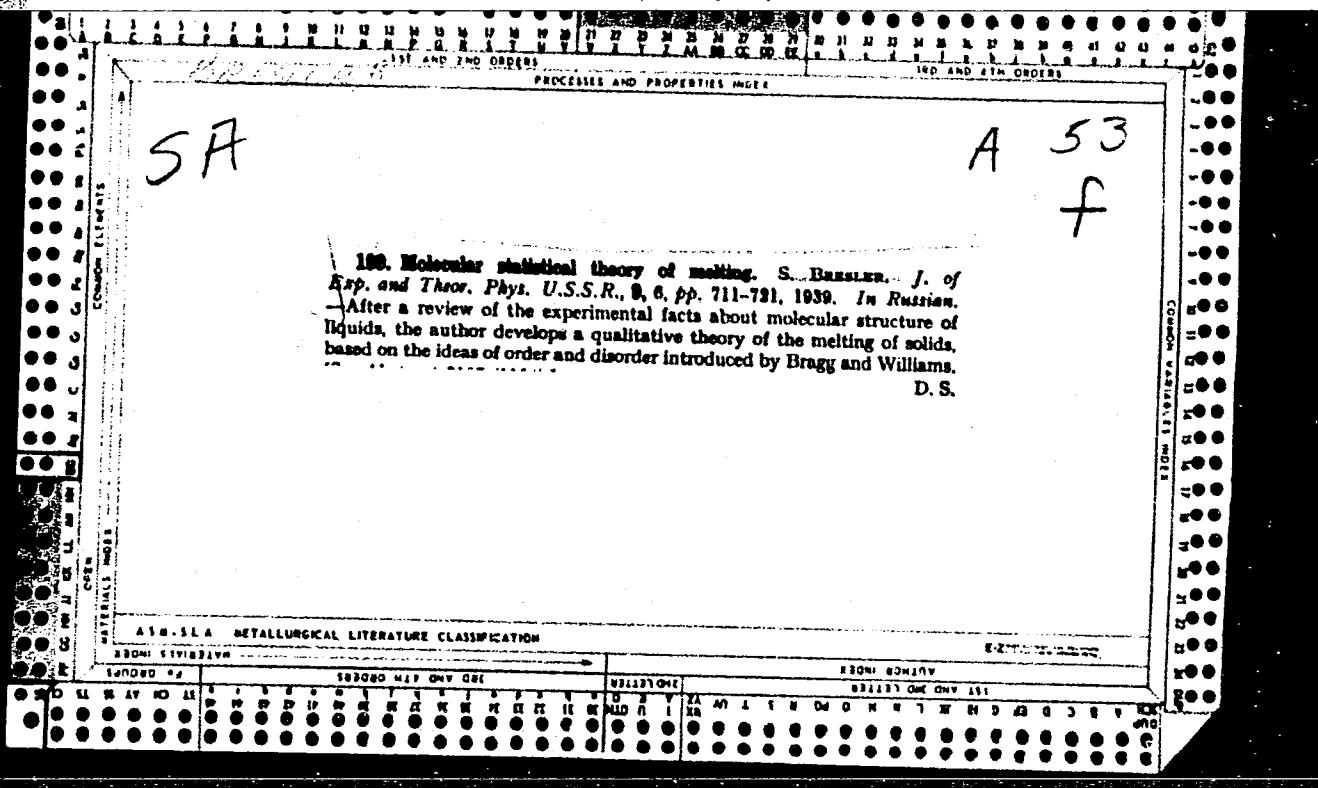
*The Structure of the Surface Layer of a Liquid (Mercury). V. P. Berdennikov, R. E. Bresler, I. Zelmanov, and P. A. Shtrauf (Zur. Fisich. Khim. (J. Phys. Chem.), 1934, 8, (5), 584-590; C. Abs., 1935, 20, 7759).—[In Russian.] (V. Physikal. Z., Norwissch., 1933, 8, 873.) In the case of mercury, the surface layer, as determined from electronograms, is quasi-solid. With the hexagonal axes of the crystallites normal to the surface.—N. B. V.

ASH-31A METALLURGICAL LITERATURE CLASSIFICATION

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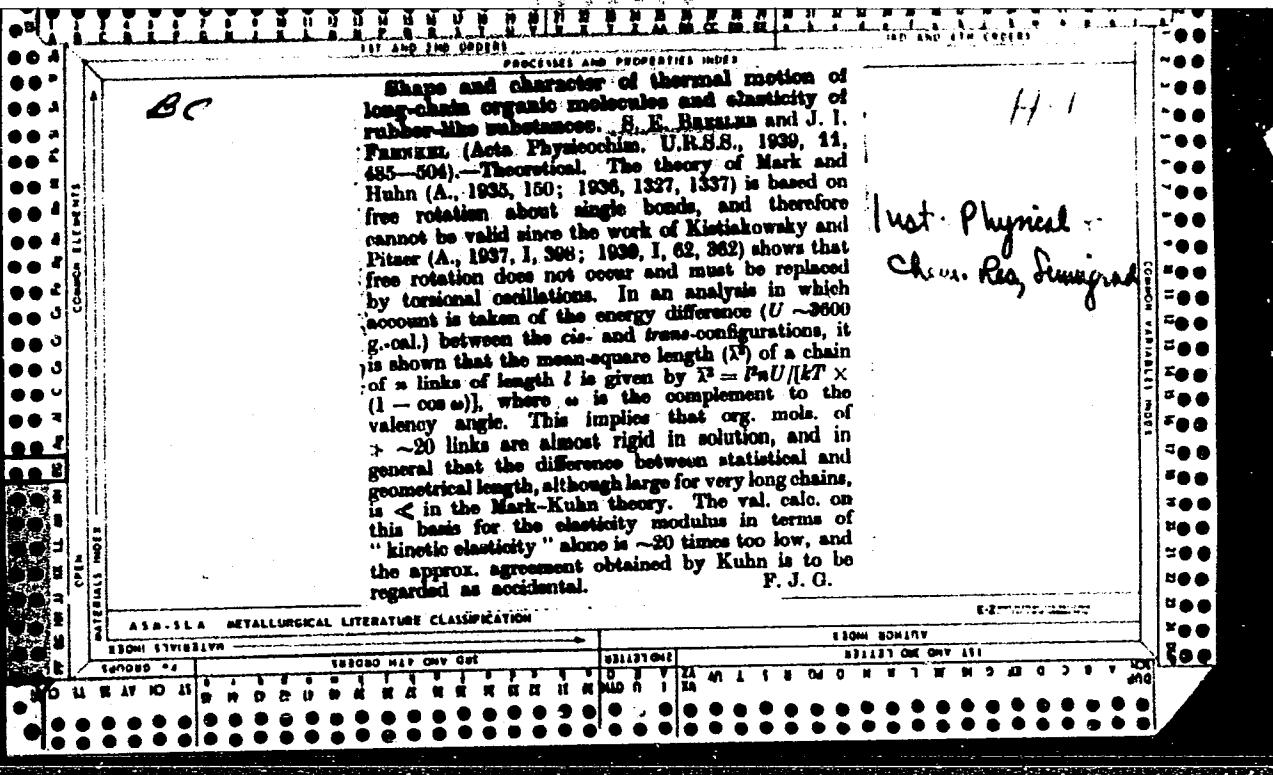




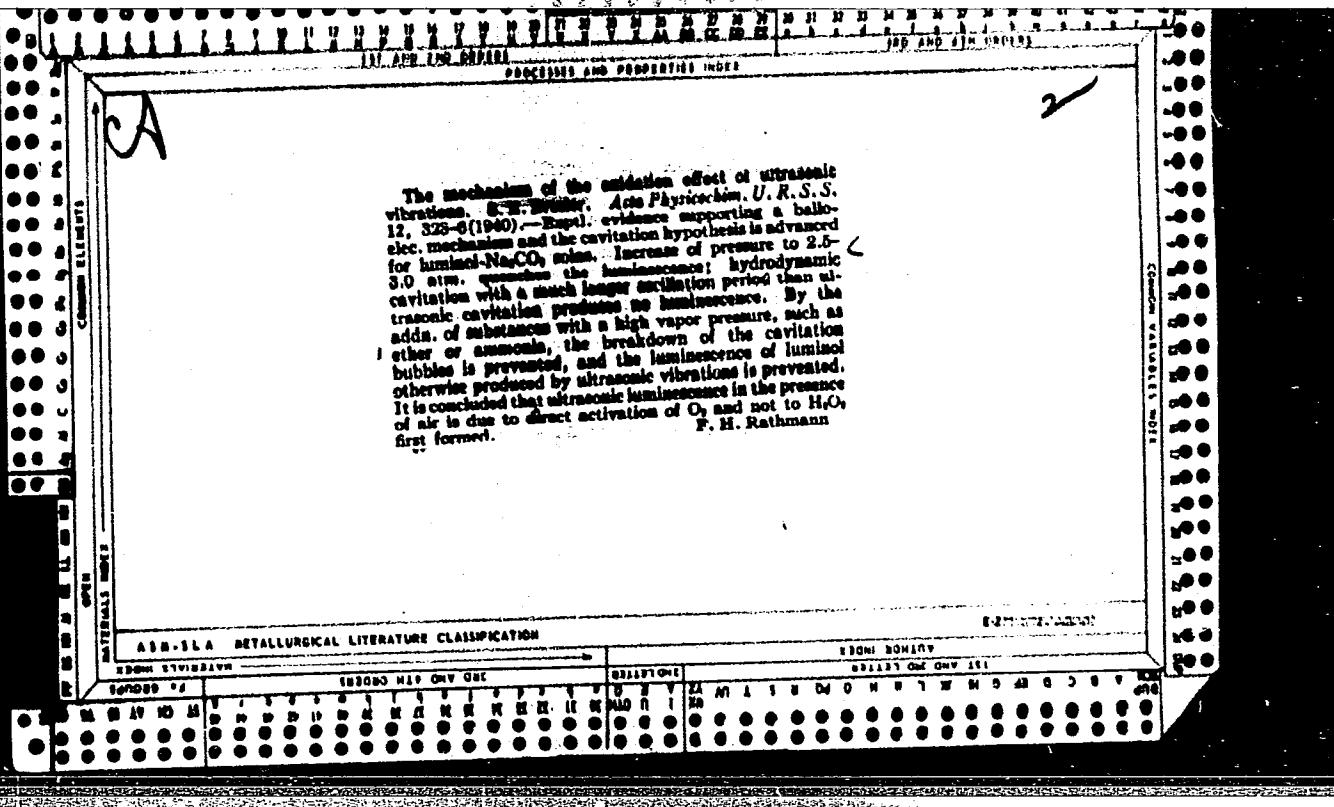
CLASSIFICATION ELEMENTS	1ST AND 2ND ORDERS		3RD AND 4TH ORDERS
	PROPERTIES AND PROBLEMS	2	
<p>The discussion of thermal motion of long organic chains with reference to the elastic properties of rubber. B. E. Brooks and Ya. I. Frankel. <i>J. Appl. Phys.</i> (U.S.S.R.) 8, 8, 1004-1106 (1939).—A crit. discussion is given of the theories of Mark (C. A. 31, 5889) and of Kuhn (C. A. 30, 7041¹). These theories are considered incorrect, since they derive the elastic properties of rubber from thermal motion alone under assumption of free rotation in long org. chains. From the latest expts. on sp. heat of ethane and propane by Pitser (C. A. 31, 4883²) and by Kastekowsky, et al. (C. A. 33, 1315³, 5373⁴) it follows that free rotation does not exist, as it is opposed by the repulsive forces between H atoms and between CH₃ radicals. The trans configuration is stable and corresponds to the min. of potential energy. Change from trans to cis state involves expenditure of energy. On the basis of these findings a new theory is developed, in which the energy of the chain is regarded as a function of the deformation angle: $U = (U_0/2)(1 - \cos \alpha)$, where α is the angle between planes passing through two neighboring links. The term $1 - \cos \alpha$ is a function of temp.: (I) $\delta = 1 - \cos \alpha = 1 - [((1 + e^{-U_0/RT})/(1 - e^{-U_0/RT})) - (2\pi T/U_0)]/2\pi^2/U_0$. Although the angle is small (close to 60°), the total deformation is equal to the sum of deformations of all links. Hence the av. statistical length of the chain (the distance between its two ends) is much smaller than its true geometrical length. For a 2-dimensional mol. it is shown with the aid of equation I that the statistical length is a function of temp.: (II) $A^2 = P^2n/\delta(1 - \cos \alpha) = P\delta U_0/2T(1 - \cos \alpha)$, where U_0 is equal to 0.6 T at ordinary temp. (Kastekowsky, et al.). α is the angle complementary to the angle between the valences of the carbon atom. It is assumed const. during the deformation process. n = no. of links in the chain; l = length of each link. For a two-dimensional model of a mol. equation (II) is simplified: (III) $A^2 = P^2n/6$, which at very high temps., when the deformation angle is small, reduces to: (IV) $A^2 = Pn$. The equation IV is very close to the expressions of Mark and of Kuhn. At very low temps. δ is less than unity, and for small n, $\delta < 1$, hence: (V) $A^2 \approx Pn^2$. This means that at low T⁵ the chain does not bend (is in the trans state), as found by Kastekowsky, et al. Thus the new theory gives correct results for the limiting cases. Equation II was applied to mols. in soln. The calcd. A is much larger than that obtained by Kuhn. This means that for aliphatic acids, amines and other mols. with n not greater than 20 the deformation is slight. An investigation by means of x-rays is thought desirable. Equation II was also compared with the Staudinger viscosity equation: (VI) $P\delta U_0/2T(1 - \cos \alpha) = A^2 k T \ln \lim[(n/n_0) - 1](1/C)\alpha - \delta$. It follows that the coeff. $k \sim 1/T$, whereas according to Kuhn's theory it should be const. Exptl. proof is thought necessary. Finally, the elastic properties of rubber were studied. When n is large the deformation is considerable, although less than that predicted by Mark or by Kuhn. If thermal motion alone is taken into ac-</p>			
Inst. Physical & Chemical Res. and Leningrad Physico-Technical Inst.			
ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION			
SECOND DIVISION		THIRD DIVISION	
SUBDIVISION	SECOND SUBDIVISION	THIRD SUBDIVISION	FOURTH SUBDIVISION
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count; then the theory of Brownian movement, together with the equation of Smoluchowsky for the diffusion of a particle, gives for the modulus of elasticity: $E = f/\Delta = \delta T(1 - \cos \omega)/m^2 = (1/m^2)[\delta(1 - \cos \omega)/U_1]^{NT}$. This equation is different from that of Kuhn and of Mark and gives much smaller E values. However, the expl. E value is larger, which leads to the conclusion that thermal (Brownian) movement alone is not sufficient for deriving the elastic properties of rubber. Other forces play essential roles in the deformation process; namely: the interaction of mols; and displacement of their centers of gravity. Also the chem. forces between the org. chains and the S atoms binding them in vulcanized rubber.

G. B. Shapiro



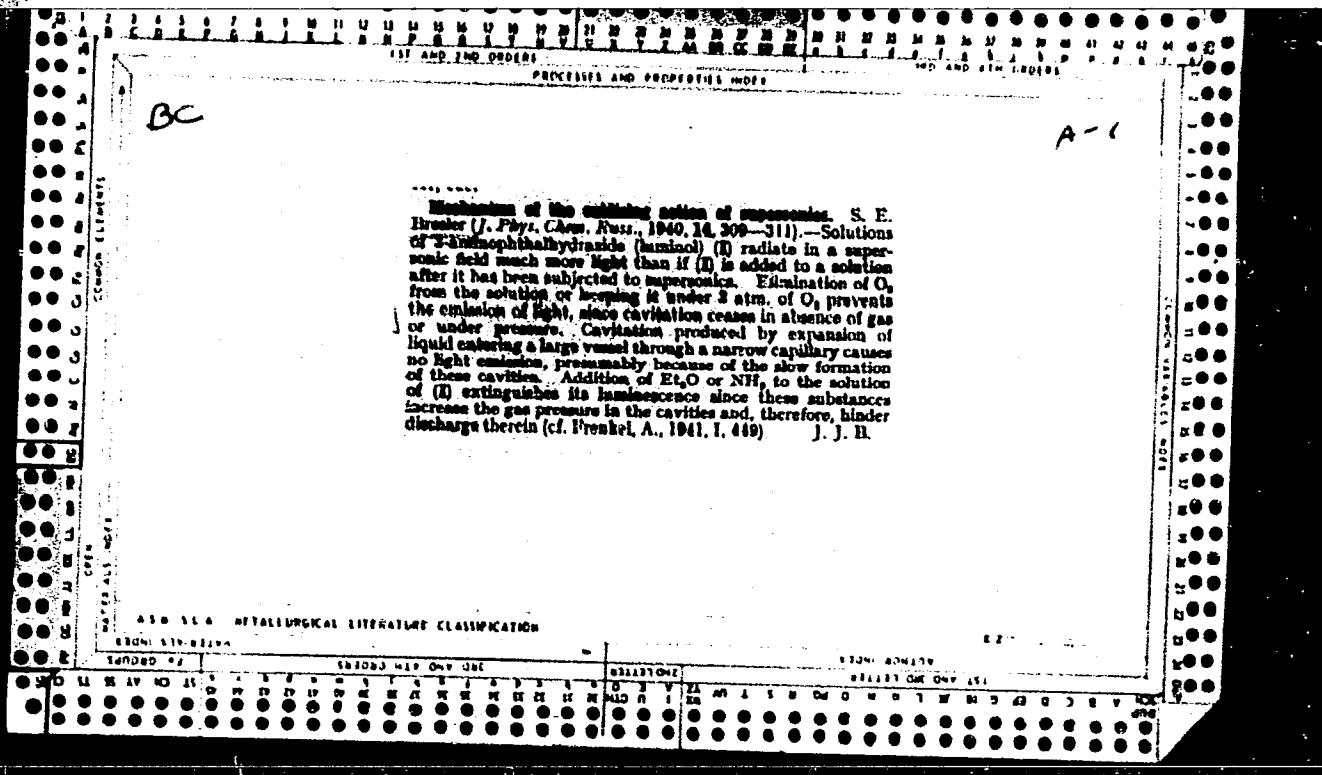
Viscosity of liquid methane and deuteriomethane.
H. Becker and A. Landerman, *J. Exptl. Theor. Phys.*, (U. S. S. R.), 10, 260-1 (1940).—According to the theory of association of liquids proposed by Frenkel and Lyman, the viscosities of liquids with the same mol. forms should be proportional to square roots of their mol. wts. Thus with liquid methane and deuteriomethane the viscosity ratio should be equal to $\sqrt{2}/\sqrt{18} = 1.110$. The viscosities of these 2 compds. were measured at the temp. of liquid O (90.1 K.) and were found to be equal to: $\eta_{\text{CH}_4} = 2.10 \times 10^{-8} = 0.01 \times 10^{-8}$ poise and $\eta_{\text{DCH}_3} = 2.34 \times 10^{-8} = 0.01 \times 10^{-8}$ poise with the ratio $\eta_{\text{DCH}_3}/\eta_{\text{CH}_4} = 1.13 \pm 0.006$. This is in very good agreement with the theoretical prediction.

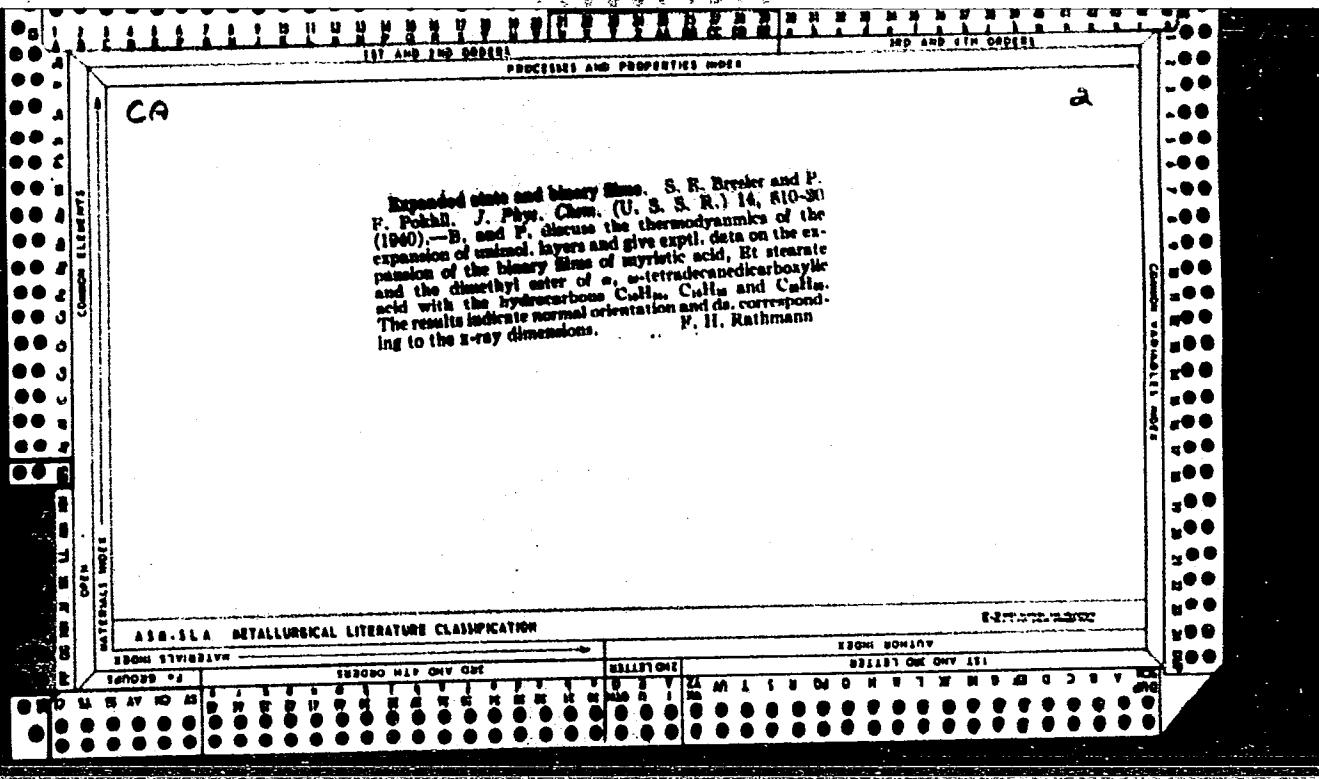


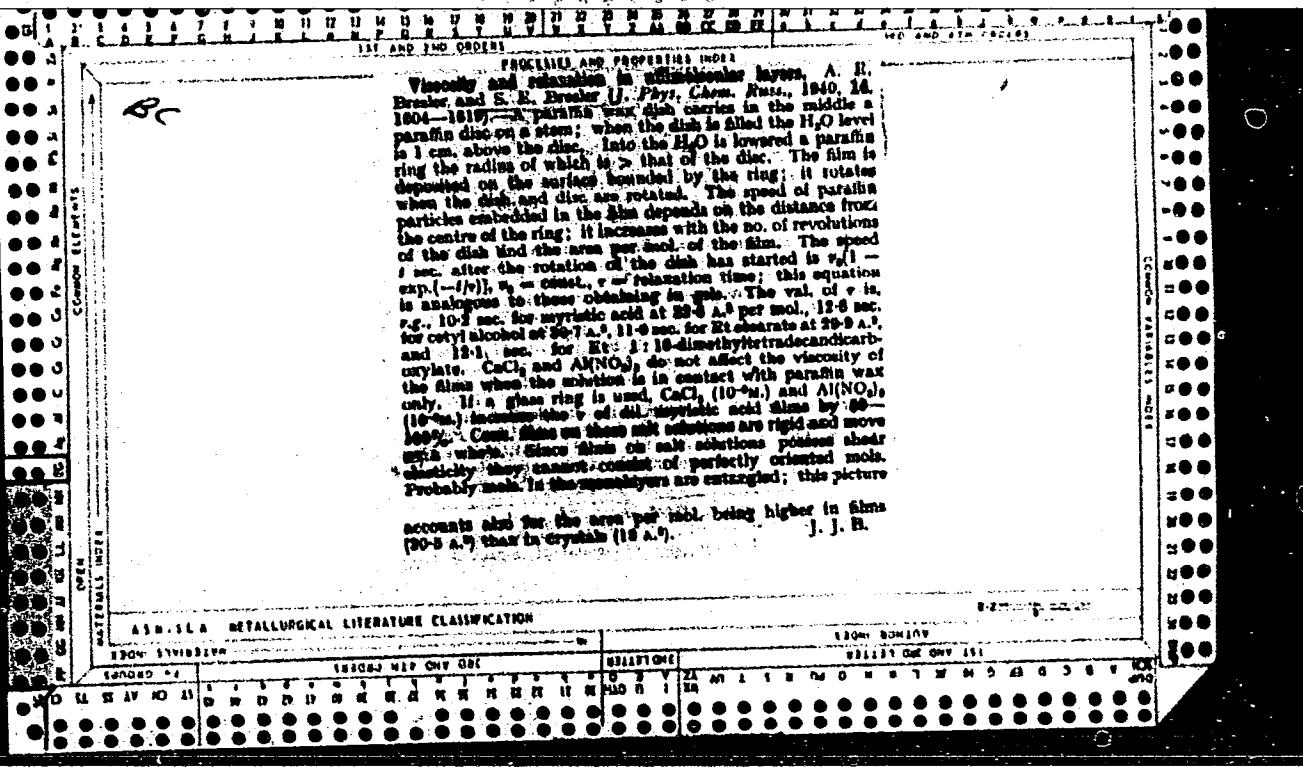
Беседка

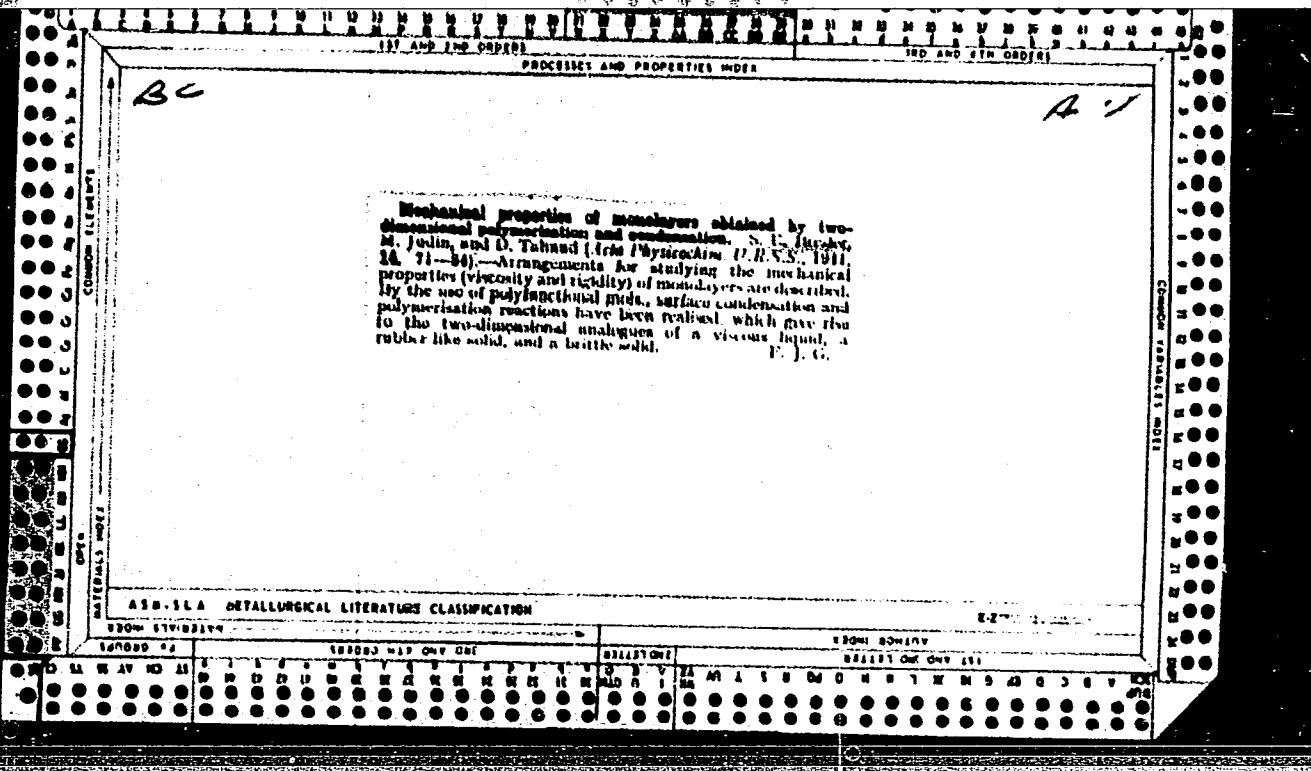
Adsorption of acetylene on selective adsorbents. S. Bresler (*Acta Physicochim. U.R.S.S.*, 1940, **16**, 783-785). The possibility of selective chemisorption of C_2H_2 on ketones of high mol. wt. has been investigated. α - C_13H_{12} O(Me), if ppd. from an alkaline medium, readily takes up $1\frac{1}{2}-1.8$ mols. of C_2H_2 per mol., the sorption being largely reversible. The activity of the adsorbent slowly decreases on keeping, and more rapidly on warming. Specimens ppd. from acid media have only a very low activity, suggesting that the active adsorbents are in the enol form, and that the loss of activity on keeping is due to a change to the keto-form. F. I. G.

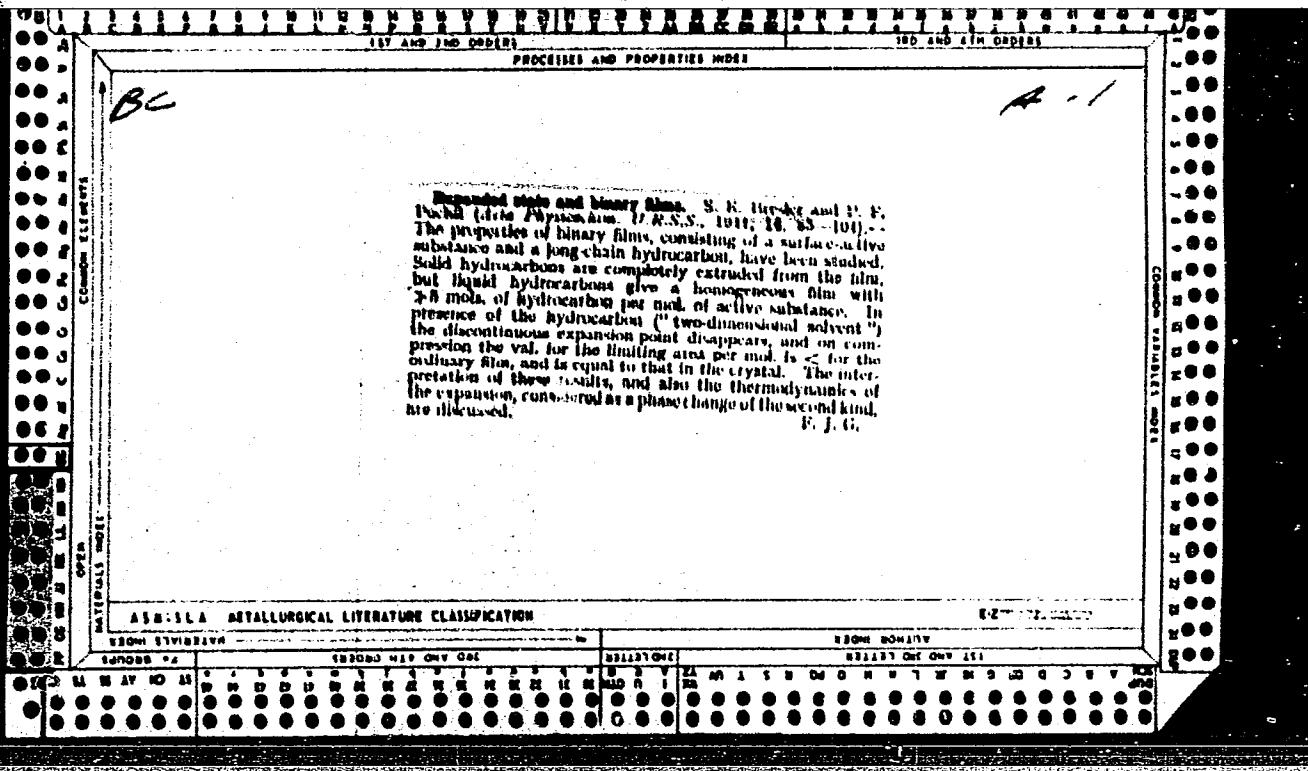
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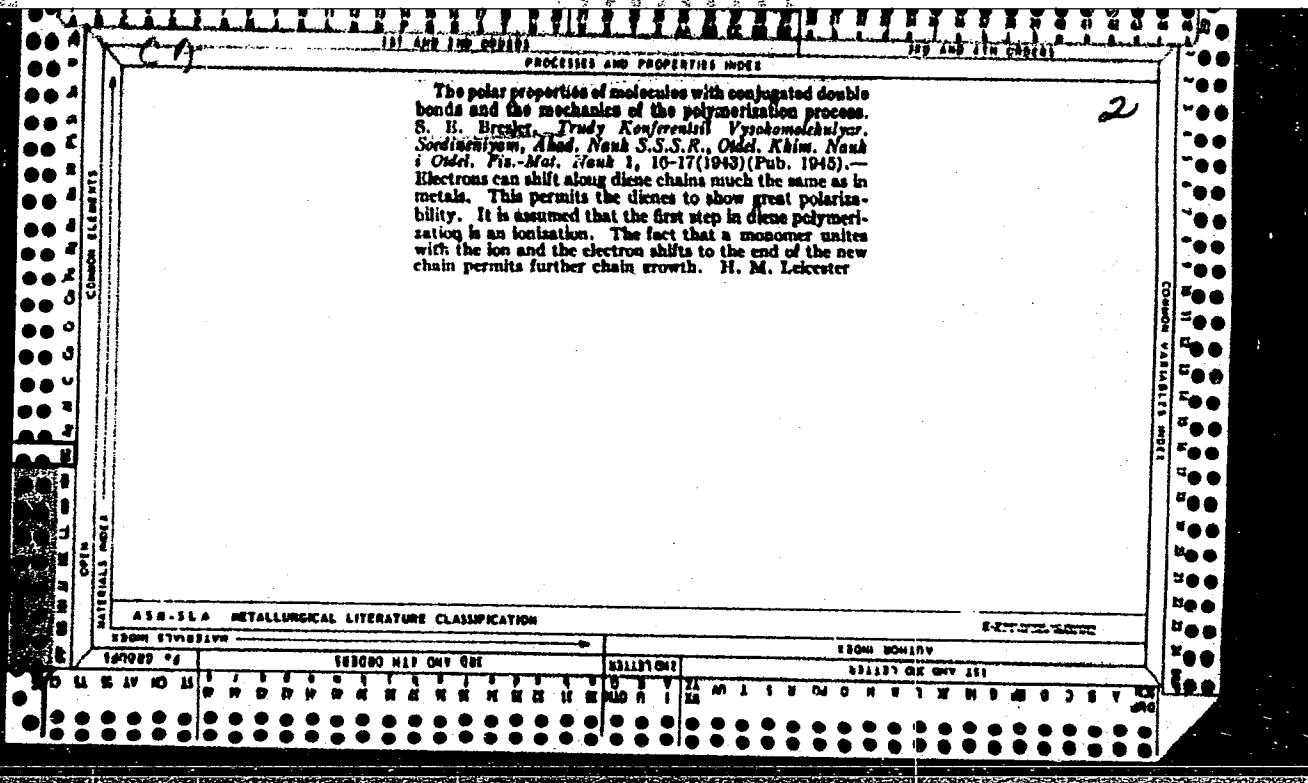












1A

2B

Vulcanization of cellulose materials. P. V. Afanasev and S. B. Breiter. *Trudy Konferentsii Vysokomolekulyarnykh Soedineniy, Akad. Nauk S.S.R., Odzsl. Akad. Nauk SSSR, Akad. Nauk S.S.R., Odzsl. Akad. Nauk SSSR, No. 2, 120-9(1944)(Publ. 1948).—Alkylation of cellulose by compds. of the type $\text{ROCH}_2\text{NR}_2\text{Cl}$ (I) is due to formation of positively charged ions of $-\text{O}-\text{CH}_2$ which are stabilized by occurrence of resonance forms. If aromatic rings or chains of conjugated double bonds are introduced, the ions are more stable and can react with OH groups in cellulose under milder conditions than can I itself. Such compds. are obtained by condensing pyrline-HCl with the bis(chloromethyl) ether of diethylene glycol, bis(2-chloroethyl) ether, the tris(chloromethyl) ether of glycerol and the bis(chloromethyl) ether of decamethylene glycol. Viscose fibers are soaked in aq. solns. of these compds. for 10 hrs. and then heated at 120° for 20 min. Vulcanization occurs and the products swell less in solvents than the original fiber. Partly hydrolyzed polyvinyl acetate and polyvinyl acel undergo the same reaction.* H. M. Lester

BRESIER, S. Ye.

Comptes Read. Acad. Sci. URSS, 1944, 43, 310-314 (cont.)

to an orderly packing of the folded polypeptide chain.

Inst. of Biochemistry, Phys, Tech .Inst. AS USSR

CA

New method of conservation of liquefied gases. M. A. Blat, S. B. Dregier, and Yu. N. Ryabinin. *J. Tech. Phys. (U.S.S.R.)* 15, 916-23 (1945).—Kistler's aerogel, a nearly transparent colloidal SiO_2 , weighing about 100 g./l., with pore dimensions of the order of 8×10^{-6} cm., was used as heat-insulating material for jackets for liquid O₂ and N₂. The aerogel vol. consists of about 5% SiO_2 and about 95% air. Pressures of the order of 1 atm. compress the powder by about 10%. Under atm. pressure, at 347°, the heat cond. is about 1.80×10^{-3} cal/cm. hr. °C. Heat cond. measurements under reduced pressures were made in an app. consisting of two concentric hollow steel spheres. The space between the spheres, containing the aerogel, was connected with a vacuum pump and two manometers, one near and other away from the pump, to insure correct determination of the pressure, between 2×10^{-3} and 20 mm. Hg. The outer sphere was kept at 115°, the inner sphere was at the temp. of the liquefied gas contained in it, -195.8°. From the amt. of liquefied gas evapd. in 1 hr., the results were plotted in terms of $\log \frac{P_1}{P_2} = \log \rho_1 - \log \rho_2$ (pressure). Without aerogel, $\log \frac{P_1}{P_2}$ depends mainly on the condition of the walls of the spheres facing each other, ρ_2 being considerably less than the walls polished. With aerogel, no such difference is observed. Examples for aerogel-insulated vessels: Space between spheres 1 cm., $\log \frac{P_1}{P_2} = -2, 0, 2, 0$; $\rho_2 =$ about 3, 12, 25; without aerogel, walls polished, $\rho_2 =$ about 18, 32, 34; space between spheres 3 cm., $\log \frac{P_1}{P_2} = -2, 0, 2, 0$; ρ_2 = about 3, without aerogel, walls polished, $\rho_2 =$ 11, 16, 37.

Only at lowest pressures, below 10^{-2} mm. Hg, are the values with and without aerogel more or less the same; above 10^{-2} mm., the insulating qualities of aerogel are striking. Its convenience lies in the fact that it requires only a moderate vacuum. Heat conductivity of the aerogel obtained in vessels of varying dimensions coincides satisfactorily. Owing to absorption, the aerogel reduces the gas pressure obtained by pumping. Tests of adsorption of air were made between 100 and 10 mm. Hg, for example, $p_1 = 20, 40, 100, 200$, and mm. Hg, absorption A_1 resp., 210, 275, 310, 365, and 415 cc. g. The plot of A_1/p_1 against p_1 is a straight line. A container made of stainless steel, with thickness 0.25 mm., cubic shape, capacity 5 L., thickness of insulation 30 mm., was tested and found satisfactory. Best results are obtained when the air is replaced by CO_2 ; this eliminates the necessity of pumping it out.

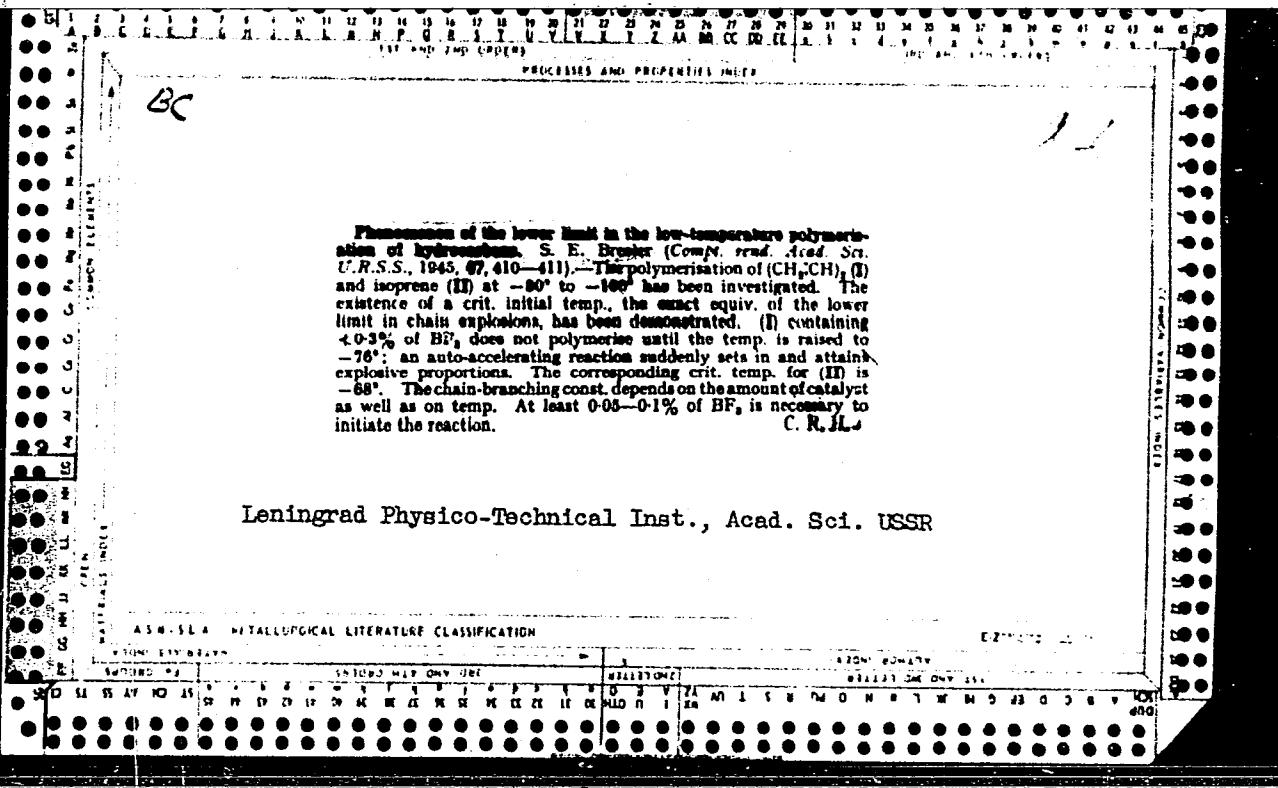
Inst. Chemical Physics, AS USSR

ASA-SEA METALLURGICAL LITERATURE CLASSIFICATION

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APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"



1ST AND 2ND ORDERS		3RD AND 4TH ORDERS		PROCESSES AND PROPERTIES INDEX	
<p><i>CA</i></p> <p>Enzymic synthesis of polypeptides at high pressures. S. K. Breker and M. V. Olikina. <i>Biofizika</i> 12, 380-403 (1947); cf. C.A. 41, 0004g.—The enzymes involved in protein metabolism usually act proteolytically when employed in the lab. However, at a pressure of 6000 atm., trypsin, papain, chymotrypsin, and pepsin exert their synthetic action, forming polypeptides and proteins from amino acids. The synthetic activity of the proteolytic enzymes at high pressures is explained by Le Chatelier's Principle. High pressures reduce the mol. vol. The expts. of Bernal (C.A. 25, 5816) are cited. Thus, in a crystal of alanine, the distance between the mols. is 5.8 Å, whereas the distance between the alanine units in alanine polypeptide is only 3.6 Å. The procedure used in the enzymic synthesis was to place the ampul with a thin capillary opening, contg. 3 ml. of the protein hydrolyzate, in a rubber sack filled with 3 ml. of the same soln. The rubber sack was placed in a thick-walled bomb surrounded by distd. water, through which the pressure was communicated by a steel piston of a hydraulic press. The rubber sack prevented contamination of the solns. with traces of heavy metals. In expts. with gelatin, a trypsin hydrolysate was employed, in a medium of 0.1 N borate buffer (pH 8.5-9.2), with a gelatin concn. of 1-4%, and the ratio of trypsin to substrate 1:30. The same gelatin soln. could be repeatedly hydrolyzed (at atm. pressure) and then resynthesized (at 6000 atm. pressure) by the action of the same sample of trypsin. The subsequent enzymic hydrolysis of the resynthesized gelatin proceeded at the same rate as the initial hydrolysis. After resynthesis, the gelatin behaved very much like the starting material; solns. gelatinized after having been cooled to room temp. However, the exact nature of the resynthesized protein (mol. wt., electrochem., and biol. properties) has not yet been elucidated. The hydrolyzed ovalbumin after resynthesis by trypsin at high pressures was also quite similar to the starting ovalbumin. The resynthesis of hemoglobin by papain yielded a product free of pigment, and resembled globulin. The result was quite different with the serum globulin resynthesized by trypsin. A gel was obtained which could not be dissolved in acids, alkalies, or in salt solns. When acted on by strong HOAc, it dispersed in the form of a turbid colloidal soln. H. Priestley</p>					
<p>Leningrad Physico-Technical Inst. Acad. Sci. USSR ASB-SLA METALLURGICAL LITERATURE CLASSIFICATION</p>					
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Bresler, S. E.

PA 20T28

USSR/Medicine - Enzymes
Medicine - Amylase

Jan 1947

"Principles of the Enzyme System Under Pressure,"
S. E. Bresler, 3 pp

"Dok Ak Nauk SSSR" Vol IV, No 2

Submitted by A. F. Ioffe, Leningrad Physical Technical Institute, Academy of Sciences of the USSR, 20 Nov 1946. Experiments carried out under pressures of 6,000 atmospheres. Concluded that under sufficiently high pressures a fermentative synthesis occurs in trypsin and amylase.

20T28

BRESLER, S. YE

PA 60T7

USSR/Chemistry - Peptides
Chemistry - Synthesis

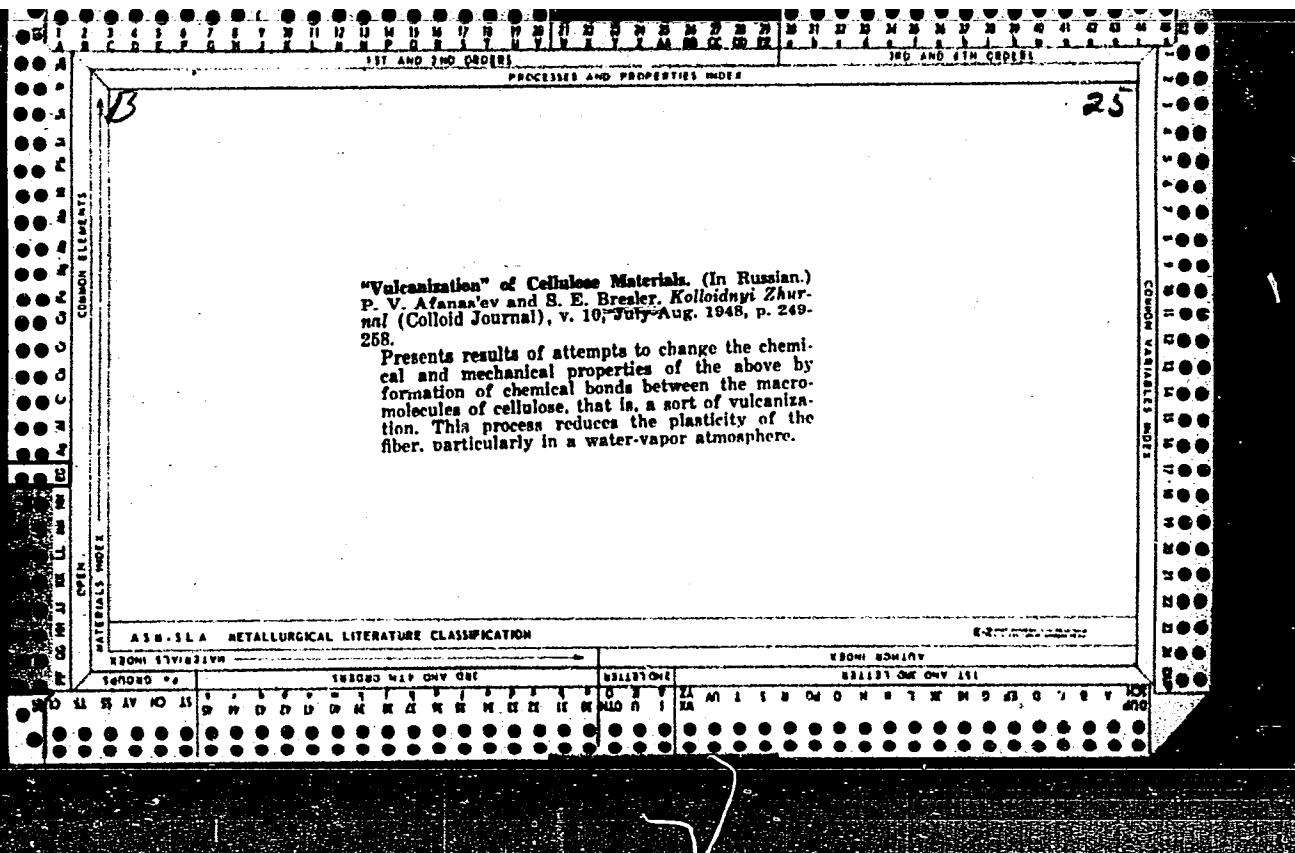
JUL 1947

"Enzyme Synthesis of Polypeptides Under Pressure,"
S. Ya. Bresler, M. V. Glikina, Phys-Tech Inst, Acad
Sci USSR, Leningrad, 4 pp

"Dok Akad Nauk SSSR, Nova Ser" Vol LVII, No 1

Describes experiments which show that proteolytic
ferments such as trypsin, pepsin, erepsin, and papain
resynthesize under a pressure of several thousand at-
mospheres. Expresses thanks to N. A. Seleznayeva who
took part in the experiments.

60T7



USSR/Chemistry -- Synthesis
Chemistry -- Albumen
Nov/Dec 48

"Problems in the Synthesis of Albumen and Starch,"
S. Ye. Bresler, Physicotech Inst., Acad Sci USSR,
16 pp

"Iz Ak Nauk SSSR, Ser Fiz" Vol XII, No 6

65-710

Concludes that fermentation reactions in
hydrolysis of albumens and starch are reversible.
Shift in equilibrium from dissolution of polymeric
molecules to synthesis is associated with variations
in volume due to changes in the peptide
(and oxide) link in the chain. Experiment re-
vealed correlation of volume variations with
synthesis phenomena for various conditions
governing reaction.

PA 25/49T10

25/49T10

The problem of synthesis of protein and starch
Proc. Tech. Inst. Acad. Sci. USSR
Leningrad Univ. Institute of Chemistry
Transl. from *Vysokomol. Soedin.*,
vol. 10, No. 10, p. 2440, 1968
by G. M. Krasil'nikov
The author wishes to thank Dr. V. A. Kabanov
for synthesis of the reducing maltose derivative
and Dr. N. N. Kostylev for help in the
synthesis of the polymer. A contribution
from the author gave after application of pressure
sure at 50° a product containing 13-15 amino acid units
with av. mol. wt. 1200-1400. Expts. on protein resyn-
thesis by means of enzymes are mentioned (cf. Bresler, C.A.
41, 6965g; 43, 704h). For such work under high pressure
the borate buffer (pH 9) appears to be the best, on the basis
of vol. changes observed during protein hydrolysis by tryp-
sin. In pure H₂O, pressure resynthesis is unsatisfactory.
Treatment of alanine in borate and soda buffer (pH 9.1)
at 5000 atm. with pancreatin led to polymer synthesis with
decline of amino-N by 23-30%. The product, purified by
adsorption on C, showed 2 fractions: alanine and the dipep-
tide. Similarly maltose was polycondensed in the presence
of malt amylase and salivary amylase; the product con-
sisted of about 8 maltose units and gave a distinct violet
color with iodine.
G. M. Krasil'nikov

USSR / Chemistry - Amino Acids
Chemistry - Proteins
May/Jun 49

"Synthesis of Proteins and Peptides Under Pressure,"
S. Ye. Bresler, M. V. Glikina, A. P. Konikov, N. A.
Selzneva, P. A. Finogenov, Molecular Dept., Physico-
tech Inst, Acad Sci USSR, Microbiol Dept., Inst of
Experimental Med, Acad Med Sci USSR, 14 pp

"Iz Akad SSSR, Ser Fiz" Vol XIII, No 3

392-406

Experiments showed that polymers regsynthesized by
authors have ~~not~~ characteristic physicochemical and
biological properties of natural proteins. A number
of important conclusions on structure of protein

52/kgt15

USSR / Chemistry - Amino Acids (Cont'd)

May/Jun 49
globule and connection of immunological and fer-
mentative activity with structure of macromolecule
can be drawn from synthesis of protein. Made
first successful steps in synthesizing amino
acids from simplest substrates. Submitted
26 Apr 49.

R.P./KOMIS

REF ID: A6561

CA

11a

Structure of globulin proteins and their interaction with the external medium. S. E. Breker. *Hishimiyu* 14, 180-9 (1940); cf. *C.A.* 36, 14283. An aq. soln. of serum albumin (1-3%) in phosphate buffer of pH 6.5 (0.03 M) on satn. with ether (6.0%), gave a lower sedimentation const. than the control, when subjected to the ultracentrifuge in Svedberg's lab., Sweden. Calen. showed an increase of 18.7% in the size of the protein mol. as a result of swelling. When 6 and 10% acetone was added to the protein soln., the mol. increase was 8 and 15.9%, resp. In the presence of urea, the sedimentation const. of serum albumin increased. The protein mol. had become heavier because 360 mols. of urea had combined with it. A soln. of serum albumin satd. with CaH_2 gave 2 sedimentation peaks, one due to the serum albumin, the other due to the bimolecular, assoc. protein. The monoprotein could not be sepd. from the diprotein by electrophoresis.
H. Priestley

Leningrad Physico-Technical Inst., Acad. Sci. USSR

ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION

CA

11 G

Immunological properties of pressure-resynthesized proteins. S. B. Bruski, A. P. Konikov, and N. A. Selezneva (Phys.-Tech. Inst., Leningrad, Acad. Sci. U.S.S.R.), *Doklady Akad. Nauk S.S.R.* 65, 521-3 (1949); cf. *C.A.* 43, 7988b.—While serum albumin loses its antigenic properties and antigen specificity upon enzymic hydrolysis, enzymic resynthesis under 6000 lb. pressure restores both properties; no new specificities arise. Hence, specificity must be ascribed not to a macromolecule but to some specific sections thereof.
G. M. Kosolapoff

BRESLER, S.Ye.; FINOGENOV, P.A.

Method of electrophoresis of proteins. Biokhimiia, Moskva 15 no.2:
145-154 Mar-Apr 1950.
(CLML 20:7)

1. Physico-Technical Institute, Academy of Sciences USSR, Leningrad.

10

New synthesis of polypeptides by condensation of amides of hydroxy acids. S. B. Bresler and N. A. Selezneva (Leningrad Phys. Tech. Inst., Acad. Sci. U.S.S.R.). *Zhur. Obshchey Khim.* (J. Gen. Chem.) 20, 350-60 (1950). AcNH₂ (40 g.) in 100 ml. abs. EtOH refluxed 20-30 min. with 11.5 g. EtONa gave a cryst. product which was taken up in more EtOAc and satd. with dry HCl, filtered, and evapd., yielding 100% $(\text{AcNH}_2)_2\text{HCl}$, m. 50°. No by-products were detected. Hence the reaction was applied to the derivs. of HO acids to form polymeric products. Lactic acid was converted by treatment of the Et ester with NH₃ into the amide which, boiled with Na in dioxane, yielded the Na deriv., $\text{MeCH}(\text{O}\text{Na})\text{CONH}_2$, m. 29°. The product (8 g.) heated in an evacuated tube 3-4 weeks to 80° gave a transparent resin, which was treated in EtOH with dry HCl, filtered, and evapd., yielding a clear resin, decomp. 105° without melting; it is sol. in H₂O, less in EtOH, insol. in Et₂O, dioxane, or Me₂CO. Condensation for 5-7 days gives a softer resin. Condensation of the free amide with metallic Na at 110° gave a dark product, aq. HCl at 33° in 22 hrs. gave 40% cleavage of the peptide links, while pancreatin gave 45% hydrolysis in 10 hrs. In both cases alanine was the end product, hence the resin was a polypeptide of polyalanine type. Adsorption on charcoal and refractometric examn. of the soln. established the polymeric nature of the product and its hydrolyzates. Polarimetric examn. showed 89% retention of the L-configuration. Mol. wt. by viscosity detns. gave 5000-8000 av. mol. wts. G. M. Kosolapoff

BRESLER, S.Ye.

Biosynthesis of protein. Usp. sovrem. biol. 30 no.1:90-112
July-Aug. 1950. (CLML 20:1)

1. Leningrad.

CA

118

Structure of procollagen macromolecules. S. E. Breiter,
P. A. Finogenov, and S. Ya. Frenkel. *Doklady Akad.*
Nauk S.S.R. 72, 635-8(1950).—Rat skin procollagen is
a monodisperse globulin as shown by ultracentrifugal
sedimentation; its diffusion coeff. at 20° in H₂O is 2.24 ×
10⁻⁷ cm.²/sec. at 0.3-0.5% concn. Its mol. wt. is estd. as
70,000 and the shape of the mol. estd. from the mol. fric-
tion coeff. is a cylinder with 20:1 axis ratio with the length
about 380 Å. Ultracentrifugal study in 0.22 M NaCl
gave a triply peaked sedimentation curve, the 2 new peaks
being caused by more rapidly moving particles, apparently
caused by lengthwise assoc., as well as axial assoc.
G. M. Kosolapoff

BRESLER, S.E.

From the Russian for Dr. Daniel Steinberg
Doklady Akademii Nauk SSSR 75, 1: 79-82, 1950

Enzymatic reaction of transfer of phosphate from adenosinetriphosphoric acid to
ribonucleic acid
by
S. E. Bresler and E. I. Nidzian

Translated at the National Institutes of Health, Bethesda, Maryland.
Full translation available in [redacted]/M.

11A

Proteolytic enzymes are metallo-proteins. S. E. Bresler and N. A. Rotentsev (Phys.-Tech. Inst., Leningrad). Biokhimiya 16, 84-94 (1951).—Com. trypsin, or the trypsin in pancreatic juice, contains 1 atom Cr per mol. protein. Cryst. trypsin (obtained by the addn. of a large excess of

$MgSO_4$) has the Cr replaced by Mg. The enzymic activity of both trypsins (blood albumin and benzylargininamide as substrates) is the same. The Cr is not firmly bound, and can be removed by dialysis. Purified, but uncrystd, com. trypsin is completely inhibited by the following org. metal complex-forming compds., in 0.002-0.005 M concns.: 8-quinolinol, formaldoxine, Na phenylthiocarbamate, thioglycolic acid, mercaptobenzothiazole, nitroso-2-naphthol, nitroso-R-salt, and diphenylcarbazide. None of these poisons which so effectively paralyze the com. enzyme have the least effect on crystd. (Mg -contg.) trypsin. On dialyzing com. trypsin, about 95% of the proteolytic activity is lost. The activity can be largely restored by the addn. of 0.0005 M Cr or 0.001 M Mg. Even the undialyzed enzymes can be activated for proteolysis by adding to the buffer solns. traces of Cr or Mg. More effective proteolysis and synthesis by proteolytic enzymes under high pressures is achieved in the presence of these metal activators. In chymotrypsin, the active metal is Mg, which is held rather firmly, and is not removed by dialysis. Chymotrypsin activity is inhibited by 0.03 M quinalizarin. It is suggested that some digestive disorders in man and in animals occur because of the scarcity of the metals entering into the compn. of digestive enzymes, or because of a metabolic disturbance of these metals in the organism. The healing effect of some mineral waters in digestive disorders is ascribed to the presence in the water of trace elements necessary to the organism.
H. Priestley

1951

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

BRESLER, S.Ye.; PAVLOVA, S.A.; FINOGENOV, P.A.

Diffusion of polymers in solutions. Zhur.Tekh. Fiz. 21,1061-5 '51.
(CA 47 no.17:8467 '53)
(MLRA 4:9)

1. Leningrad Phys.-Tech.Inst.

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

CIA
11A

Resynthesis of biologically active insulin. S. E. Brooker,
M. V. Glikina, and A. M. Tongur. *Doklady Akad. Nauk S.S.R.* **78**, 543-5 (1951).—Cryst. insulin as a 0.6% soln. in 0.2M borate buffer at pH 8.8 was used as starting material. This was hydrolyzed by 0.0012% trypsin and 0.0003% chymotrypsin for 4-5 hrs. to the extent of about 5% which is max. for these enzymes. Since the enzymes are inactivated under pressure, they were stabilized by 20% glucose. Pressure resynthesis, measured by decrease of free amino groups, reached 96%. To prevent renewed hydrolysis by the enzymes on release of pressure the mixt. was adjusted immediately to pH 2.5 and frozen in liquid air. The resynthesized insulin, in contrast with the hydrolysate, showed typical insulin sedimentation consts.; this indicated reestablishment of its macromol. from proteolytic fragments of low mol. wt. The bio. activity was re-established to the extent of some 10%. The hydrolysate used as starting material for resynthesis was inactive.
G. M. Kosolapoff

NW/ 7691

CII RADIOACTIVE ELEMENTS. S. E. Bresler. Moscow-Leningrad, Gostekhizdat, 1952. 735p. (In Russian) (Book on display at Geneva Conference)

A monograph on the chemistry of radioactive substances and radiochemical methods of investigation. Radioactive series; analysis of the processes of concentration, separation and adsorption of radioactive substances. Fundamentals of the chemistry of radionuclides and artificial radioactivity and of nuclear reactions. Methods of producing artificial radionuclides and their compounds. Application of labelled atoms in chemical, biological, medical and geological research; bibliography after each chapter. (publisher's note)

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BRESLER, S. Ye.

Brit Ab RIII
July 1953

General Organs +
Constituents.
Comparative Physiology

✓ Resynthesis of protein under pressure. S. E. Bresler, M. V. Glikina, N. A. Sclernevaya, and P. A. Finegenov. *Biochimia*, 1952, 17, 44-55. Various proteins were treated with crystalline proteases. After hydrolysis had occurred the mixture was exposed to a pressure of 6000 atm. and the resynthesis of the protein studied. It is found that with mixtures of different substrates, e.g., ovalbumin, serum albumin, the amount of protein resynthesised was greatly reduced, although each of the proteins separately could be resynthesised. Resynthesis takes place not in steps but rapidly, and it is not possible to isolate any intermediary substances between the products of hydrolysis and the resynthesised protein.

(S) Chern

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4-26-54

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

BRESLER, S. YE., PAVLOVA, S. A., FINOGENOV, P. A., CHNUTOV, K. V.

Polymers and Polymerization

Remarks on the paper "Diffusion of polymers in solutions" Zhur. fiz. khim. 26 No. 3, '52.

9. Monthly List of Russian Accessions, Library of Congress, September 195~~6~~? Uncl.

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

USSR/Chemistry, Biological - Proteins 11 JUN 52

"Crystallization of Resynthesized Protein," S. Ye.
Bresler, N. A. Selezneva

"Dok Ak Nauk SSSR" Vol LXXXIV, No 5, pp 1013-1015

Authors showed formerly that enzymatic resynthesis under pressure of products of deep fission of proteins yielded artificial substances of globular structure which exhibited a biol activity (antigenic, enzymatic, and hormonal) typical for native proteins. In the expts described now, equine serum albumin was split by trypsin and chymotrypsin. Upon addn of glucose serving to protect the enzymes against fission,

223T22

the protein was resynthesized under pressure. Investigation in an ultracentrifuge showed that the resynthesized protein showed a mol wt dispersion of 22% in comparison with the initial albumin. This dispersion explains the lower biol activity per unit of wt exhibited by resynthesized proteins. The product of resynthesis crystd with great facility, however.

223T22

BRESLER, S. YE.

"APPROVED FOR RELEASE: 06/09/2000 CIA-RDP86-00513R000306910006-6

BRESLER, S.E.; KORSHAK, V.V.; PAVLOVA, S.A.; FINOGENOV, P.A.

Experimental study of the molecular-weight distribution function of polyamides. Doklady Akad.Nauk S.S.R. 87, 961-4 '52. (MLRA 5:12) (CA 47 no.14:6738 '53)

1. Inst. High-Mol. Compds., Acad. Sci. U.S.S.R., Moscow.

APPROVED FOR RELEASE: 06/09/2000 CIA-RDP86-00513R000306910006-6"

BRESLER, S.YE.

"Concerning the Article 'Synthesis of Proteins' by S.Ye. Bresler
V.P. Korotkoruchko, Kiev, reviewer)
Biokhim, vol. 18, no. 1, pp 130-136, Jan/Feb 1953

Review is an exhaustive criticism of an article which appeared in Voprosy Filozofii,
no.3, 1951, pp 82-94. The reviewer accuses Bresler of deviating from the theories of
dialectical materialism and of presenting theories which are in direct contradiction
to his own (Bresler's) experimental data. He quotes Bresler as denying the
existence of polypeptides in living tissue and accuses him of disregarding the effects
of environment on the synthesis of proteins. His general conclusion is that Bresler
has brought considerable confusion to the accepted ideas on the biological synthesis
of proteins.

257T3

Enzymic synthesis of the oside bonds under pressure
S. E. Bresler and S. Ya. Frentsel (Inst. of High Mol.
Compounds, Acad. U.S.S.R., Leningrad). *Biochemistry*
19, 648-651 (1973). Under pressure of 0.5 thousand atm,
it is possible to convert the amylolytic reaction into a
process of synthesis. α-1,4-Glucoside was thus synthesized
by two methods described. Unlike the case of protein
synthesis, no high mol. products could be obtained. Be-
cause of the loss of dextrinogenic activity, amylases are
ineffective as catalysts in the interpolymerization of dextrin
B. S. Levine

USSR.

The theory of nonequilibrium chromatography. S. E. Bresler and Ya. S. Ulyanov. Zhar. Tekh. Fiz. 23, 1443-51 (1953).—A differential equation is written for the condition that the adsorption on the grains of the adsorbent is not an equal. value, but varies by a time t . This equation is $v(\partial v/\partial t) - (\partial c/\partial t) - (w/v_b) - (\partial c/\partial z) = 0$, where w is the velocity of the liquid in ml./sec., v the cross section of the column, k a const. (the sum of the adsorption const. and the coeff. of porosity), and c the concn. This equation was solved by means of Laplace transform both for the case of frontal and of elutri chromatography. S. Pakswar.

H. 82

✓ 5879. Investigation by means of an ultracentrifuge of the molecular weight distribution of polymers.
S. E. BROWNE and S. V. FRENKEL. Ztsur. Tech. Phys., 1953, 23, 1602-20; In English: Russ. Chem. Technol., 1957, 80, 487-600. A method is given for studying molecular weight distribution of linear polymers in general. It involves fractionating the polymer into a series of narrow fractions and investigating each with the ultracentrifuge; plotting the distribution functions of sedimentation constants for each fraction and subsequent summation of curves to build up the distribution function of the sedimentation constant of the whole polymer; discovery of a general functional relation between sedimentation constants and molecular weights for a given series of polymer homologues; and construction of the distribution function of molecular weights of the polymer.

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BRESLER, S. E.

U.S.S.R.

The molecular-weight distribution of polymers. S. E. Bresler, I. Ya. Poddubnyi, and N. Ya. Frenkel. *Vysokomol. Soedin.*, Vol. 23, No. 1, p. 1521-40 (1953).—Three samples of rubber were selected: 2 butadiene polymers with plasticity 0.3 and 0.20 and 1 low-mol. (plasticity 0.5) butadiene-styrene copolymer (Buna-S). A mol.-wt. distribution was obtained by fractional pptn. by methanol from benzene solns. The fractions were dried in a vacuum and weighed. Their mol. wts. were detd. by osmometry, viscometry, and diffusion of light. The resulting step curve of distribution was smoothed out and compared to the curve obtained from ultracentrifuge data by the method described in the preceding abstract. The latter gives more details about the distribution curve. The calcs. of corrections, included in the method, is given in detail. S. Pakswert

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BRESLER, S. YE.

The Committee on Stalin Prizes (of the Council of Ministers USSR) in the fields of science and inventions announces that the following scientific works, popular scientific books, and textbooks have been submitted for competition for Stalin Prizes for the years 1952 and 1953. (Sovetskaya Kultura, Moscow, No. 22-40, 20 Feb - 3 Apr 1954)

<u>Name</u>	<u>Title of Work</u>	<u>Nominated by</u>
Bresler, S. Ye.	"Resynthesis of Biologically Active Proteins Under Pressure"	Institute of Biological and Medical Chemistry, Academy of Medical Science USSR
Glinkina, M. V.		

SO: W-30604, 7 July 1954

Disturbance of chemical equilibrium in surface layers
S. E. Bresler and G. V. Samsonov (Acad. Sci. U.S.S.R.).
Zhur. Fiz. Khim. Nauk. S.S.R., *Ved. Khim. Nauk* 1954,
p. 23. A study was made of the problem of the effect of
chemically interacting substances in the surface layer.
Investigations in the direction of chem. processes led to an
experiments applied to the surface of a liquid. The process
of synthesis of esters from acid and alcohols in monolayers
exists in systems consisting of a mixture of a fatty acid and
alcohol or of a combination of different organic acids. A
method of Langmuir Adams, as well as by the help of
electrolytic conductometric analysis. A series of organic
acids (fatty acids) in palmitic acid were studied, and a series of alcohols
from methanol to octadecyl was used. Unusual surface
concentrations were applied to the surface of water in the Langmuir
method. Surface tension was measured by means of a torsion
balance by using the method of breaking away of a glass plate
leaving a perimeter of 20 mm. The error involved in this
measurement was 0.25 dynes/cm. For applying the film a
0.1% soln of acid and alc in hexane was employed. This
solution was applied in the quantity of 6-6 drops from a specially
calibrated micropipet. The esters formed were recovered
and subjected to colometric analysis after addition of
phenolphthalein to produce a red color. The synthesis of ester
from a fatty acid and an alc was demonstrated and it was
shown that under conditions of monolayers reactions of
dehydration proceed in the presence of practically endless
reservoirs of water, and that the original conditions in the
surface layer displaced the chem. equil. from hydrolysis to
synthesis. Data are presented graphically. G. S. M.

BRESLER, S.Ye. (Leningrad)

Enzymatic synthesis of protein. Usp.biol.khim. 2:66-96 '54.

(MIRA 12:12)

(PROTEINS,
synthesis, fermentative)

136 ESTER SUE

✓ Investigation of the distribution function for the molecular weights of polyamides by sedimentation and ultracentrifuging. I. Determination of the distribution function from sedimentation constants, II. Molecular weight distribution of polyamides and mechanism of condensation. S. E. Bresler, V. V. Korshak, S. A. Pavlova, and P. A. Finogenov. (Izvest. Akad. Nauk SSSR, Otdel. khim. Nauk, 1954, No. 2, 344-353, 354-381).—I. Distribution functions for mol. wt. are deduced for polyamides by sedimentational (ultracentrifuge) analysis of fractions obtained by fractional precipitation of the polymer with ether from methyl alcohol. After correction for the effects of inhomogeneity of the centrifugal field and of diffusion, the sedimentation curves approximate closely to Gaussian curves and the calculation of the distribution function is simple. II. Distribution functions derived from sedimentation constants and diffusion coefficients by a fairly vigorous method do not coincide with the theoretical ones calculated on the basis of Flory's theory of the mechanism of linear condensation. An alternative theory is propounded, making allowance for exchange and degradational reactions between molecules of polymer, and of polymer and monomer. The existing methods of fractionating polymers are criticised and the state of polyamides in solution is discussed.

R. C. MURRAY

BRESLER, S. E.

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(6)

The molecular weight of plasteins. S. E. Bresler, K. S. Makarov, and S. Ya. Frenkel (*Izst. High-Mol. Compds., Acad. Sci. U.S.S.R., Leningrad*). *Biokhimiya* 19, 88-95 (1954).—Sedimentation and diffusion consts. and mol. wts. of two plasteins were detd. The plasteins proved to be polydispersed low-mol. peptides with an av. mol. wt. of about 500. The presence of high-mol. fractions in plastein compns. reported by others are explainable on the basis of secondary aggregation. Plasteins contain no heavy-mol. fractions. The formula of Gutfreund and Ogston (*C.A.* 43, 6259) yielded correct sedimentation consts. for low-mol. polypeptides. Calcn. of mol. wt. on the basis of distribution in the diffusion layer close to the bottom of the tube can be made with the aid of the barometric Boltzmann formula.
B. S. Levine

10-15-54

mm

BRESIER, S. Ye., PRYADILLOVA, V. I. and KHAINMAN, V. Ya.

"Investigation of the Mechanism of Rubber Vulcanisation aided by Radioactive Sulphur. I." Zhur. Tekhn. Fiz., 1954, 24, p. 577-98

Thin rubber discs were coated on the under surface with sulphur, isotope 35, and radiation measured on the upper surface by a gieger counter. There was no cross-linking in natural rubber, using 0.01 to 0.02% S without an accelerator, the sulphur concentration becoming equal on both surfaces. Treatment with thiuram type accelerator or mercaptobenzthiazole, 0.3 to 0.5%, resulted in a low concentration on the upper surface. Combined and free sulphur were directly proportional, corresponding to a first order equation. Accelerators with 3% sulphur showed secondary cross-linkage due to reverse diffusion. Radiation from the upper surface reached a maximum and then dropped by 25%. This indicated the sulphur concentration on the lower surface to be temporarily lower than that on the upper. Buna has only primary addition whatever the sulphur concentration. The ~~first~~ diffusion constant shows the sulphur molecule to be S_8 . Only two thiuram sulphurs, those of the disulphide group, take part in the exchange reaction, S^{35} -thiuram. These form a thermodynamically unstable active sulphur biradical, which breaks down a S_8 ring to further biradicals, which then bond hydrocarbons already activated by primary addition. Each polysulphide bond has a length of 10 to 20 sulphur atoms. The scission energy (2,700 cal/mol) suggests the diffusion of most of the bonded rubber. Though vulcanised natural rubber has considerable diffused sulphur, a Buna vulcanisate has little, due to the small number of polysulphide bonds.

BRESLER, S. Ye., KUSHNER, V. P. and SAMINSKIY, Ye. M.

"Study of the Mechanism of Vulcanization of Rubber with the aid of Sulfur-35. II"
Zhur. Tekh. Fiz. 24, 2150-68, 1954.

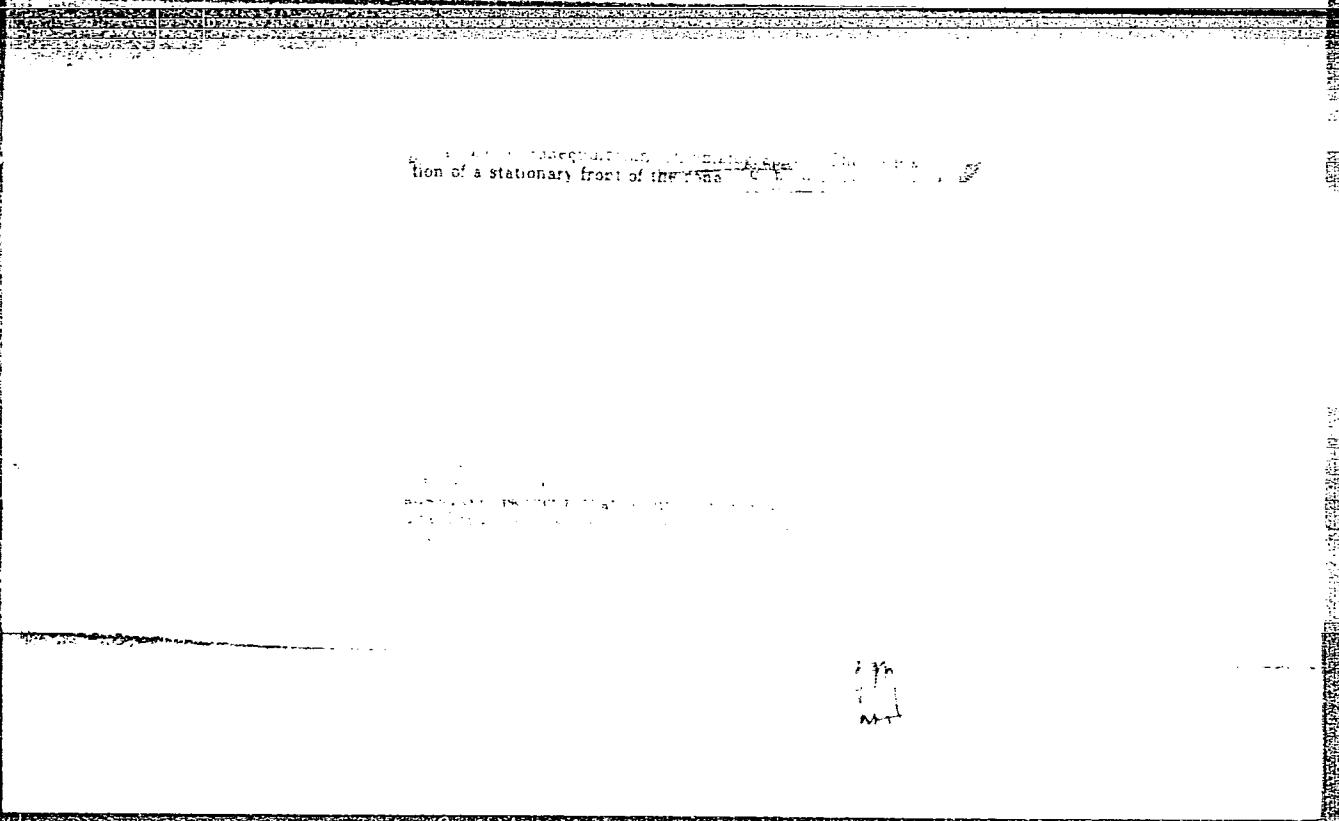
In various synthetic rubbers, and especially natural rubber, a nonlinear addn. of S³⁵ was observed at concns. >1% and at elevated temps (120-155°). The primary addn. of S was found to be partially followed by the secondary reaction of S-forming polysulfide bridges between the rubber chains. The departure from simple kinetic characteristics of the process is explained on the basis of calcns. of the diffusion coeff. and the amt. of polysulfides in the rubber. Similar measurements were also carried out for rubber vulcanized under the usual conditions (1-2% S; 130-140°). The free radical S₂-- reacts with rubber, even at room temp. The app. is described in detail.

Enzymic activity of fragments of protein of low molecular weight and the presence of the trypsin inhibitor in the fragments.

The 1% soln. was autoclaved in borate buffer at pH 7.5 at 37° for 3.5 hrs., the process being followed by centrifugation at 10,000 rpm for 15 min. in a large ultracentrifuge cell. The supernatant was then dialyzed against 0.05M Tris-HCl, pH 7.5, for 16 hrs. After dialysis, the solution was concentrated to 10 ml. by lyophilization. Ultracentrifuging was continued 6-8 hrs. at 65,000 r.p.m. after acidification to pH 3 and examination of a specimen taken from the upper part of the cell (free from protein macromolecules) showed that the products were substances of relatively low mol. wt. (2000-3000). This material had protease activity which was only reduced from that of the original enzyme by a factor of 16-20. A control run, in which whole tissue was treated similarly, also showed a weak activity in a fraction of low mol. wt. taken from the upper part of the ultracentrifuge cell; this, however, was several times weaker than that found for the autoclaved specimen. A specimen of myogen was hydrolyzed by combined action of trypsin and chymotrypsin to the extent of 5-6%, after which the product was ultracentrifuged as above. In this case the solution obtained after dialysis, treatment with 0.05M Tris-HCl, pH 7.5, and concentration to 10 ml. showed a marked reduction in protease activity. These results indicate that the fragments of protein of low molecular weight are active enzymes.

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6



APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

BRESLER, S.Ye.

Resynthesis of protein under pressure. S. B. Bresler
(Inst. Highmolecular Compounds, Acad. Sci. U.S.S.R.,
Leningrad). Biokhimiya 20, 463-6(1955).—Talwar and
Macheboeuf (C.A. 48, 6477b) and Macheboeuf, et al. (C.A.
49, 2038c) reported their inability to reproduce the results
previously reported by Bresler, et al. (C.A. 46, 5830c), re-
garding the resynthesis of proteins under pressure.¹ The
difference in the results obtained was because the French
authors probably used impure enzyme preps. A method
of enzyme purification was described and new data presented
which confirm previous assertions. B. S. Levine

CH

Bresler, S Ye.

EXCERPTA MEDICA Sec.2 Vol.10/2 Physiology, etc Feb57

572. BRESLER S. E. and RUBINA Kh. M. Biochem. Dept., 1st (Pavlov) Med. Inst., Leningrad. *Enzymatic transportation of phosphate groups from ribonucleic acid to fructose monophosphate (Russian text) BIOKHIMIJA 1955, 20/6 (740-748) Tables 4
Phosphorylated yeast RNA has been enzymatically synthetized. The substance obtained in the presence of 6-phosphofructokinase delivers the phosphate group to fructose 6-phosphate, forming fructose diphosphate. It is believed that the phosphorylated RNA is a high-energy phosphate, resembling ATP. Enzymatic participation of 6-phosphofructokinase seems to indicate that the transported phosphate group must be bound to the 5th carbon of ribose. Szabuniewicz - Gdańsk

"APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6

APPROVED FOR RELEASE: 06/09/2000

CIA-RDP86-00513R000306910006-6"

BRESLER, S.Ye.

The investigation of the reaction mechanism of polymerization by the [measurement of the] distribution of the molecular weights in the products. S.Ye. Bresler and S. Yu. Frenkel, Zhur. Tekh. Fiz. 25, 2160 (1949) [cf. U.S.A. 49, 3023].—11 polymers, which are not identified, were send into the components of various mol. wt. by an oil ultracentrifuge at 40,000 r.p.m. (100,000 $\times g$). The temp. during the runs never rose more than 0.5° and was between 15 and 18°. The diffusion coeff. was detd. in an app. where the temp. could be kept const. to $\pm 0.003^\circ$, and the viscosity was measured in a suitable viscometer. The reference value for all values is the sp. vol. of polystyrene in Me Rt ketone of 0.91 cc./g. Werner Jacobson

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BRESLER, S. Ye.

Category: USSR / Physical Chemistry - Surface phenomena. Adsorption.
Chromatography. Ion exchange.

B-13

Abs Jour: Referat Zhur-Khimiya, No 9, 1957, 30218

Author : Samsonov G. V., Bresler S. Ye., Vansheydt A. A., Kuznetsova N. N.,
Lavrent'eva S. F., Shesterikova M. P.

Inst : not given
Title : Sorption of Streptomycin by Carboxyphenol Resins

Orig Pub: Antibiotiki, 1956, 1, No 5, 42-46

Abstract: Trivalent cathions of streptomycin (Str^{3+}) are sorbed irreversibly at sulfocathicnites while with purely carboxylic cathionites (KFU and KMT) absorption capacity for Str^{3+} amounts to only 38-22% of their capacity for simple inorganic cathions (Na^+ and Ca^{2+}), evidently due to steric hindrances caused by excessively close distribution of carboxyl groups. It was found, in accord with the theoretical assumption, that the readily swelling, capable of ion-exchange

Card : 1/2

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Ion-exchange chromatography of streptomycin
mulfite. S. E. Bresky and G. V. Samsonov (Inst. H.
McL. Compos. Acad. Sci. U.S.S.R.)

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CIA-RDP86-00513R000306910006-6"

BRESSLER, S.E.
Chemistry.

~~S. B. F. SLEATOR, D.Sc.~~ USSR Physical Chemistry. Surface Phenomena. Adsorption. Chromatography. Ion Exchange.

B-13

Abs Jour : Ref Zhur - Khimiya, No 7, 1957, 22561.

Author : G. V. Samsonov, S. E. Bresler.

Author : G. V. Sam
Inst : Not given

Author : Not given
Inst :
Title : Statics and Dynamics of Streptomycin Ion Exchange with Metal
and Hydrogen Ions on Carboxyl Tars.

Orig Pub : Kolloid. Zh. 1956, 18, No 3, 337-343 (rez. angl.)

Abstract : Carboxyl cationites (CC) unlike sulfocationites sorb streptomycin (I) reversibly from solutions, but only in case when CC are used in Na^+ , K^+ , or NH_2^+ salt form. Exchange capacity of CC in relation to I depends on the degree of swelling, which creates new possibilities for increase of accessibility of ionite active centers for large ions I, in comparison to permuites (RZhKhimii, 1956, 57703). Equilibrium of the I ion exchange with metal cations conforms with B.P. Nikol'skiy's equation, if we take in account only those metal ions in cationite which can exchange with I. For one g-mole of I - 3g-mole of Na^+ are displaced and as a result of that a

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CIA-RDP86-00513R000306910006-6"

*Description and chromatography of Aureomycin G V
K. V. S. B. Rostovtsev and A. A. Kozentsev (Inst.
High-Mol. Compds., Acad. Sci. U.S.S.R., Leningrad)
Naukova Zbirka, 19, 170-3 (1956). Aureomycin-HCl (I),
from a 0.05% soln. was most strongly adsorbed by C. at*

Leningrad Chemico-Pharmaceutical Inst.

Category : USSR/Atomic and Molecular Physics - Liquids

D-8

Abs Jour : Ref Zhur - Fizika, No 1, 1957 No 970

Author : Bresler, S.Ye., Pikus, G.Ye.
Title : On the Separation of Ions by Their Mobilities.

Orig Pub : Zh. tekhn. fiziki, 1956, 26, No 1, 109-125

Abstract : Development of a phenomenological theory of the separation of ions by their mobilities, using as an example the separation of isotopes of liquid metal by electrolysis. Expressions are derived for the stationary and non-stationary distributions of the concentration of the isotopes and for the amount of isotope concentrated at the edge of the tube (for the stationary cases). The laws derived are applied to the analysis of the experimental data on the separation of Hg and Ga and to the calculation of the differences in the mobilities of the isotopes of these elements.

Card : 1/1

Bresler, S.Ye.

USSR/Chemistry of High Molecular Substances.

F

Abs Jour: Ref Zhur - Khimiya, No. 8, 1957, 27064.

Author : Bresler, S.Ye., Dolgoplesh, B.A.;
Inst : Krol', V.A., Frenkel', S.Ya.

Title : Reactions of Free Radicals in Solutions. V.
Destruction of Polymer Molecules under Influence
of Free Radicals.

Orig Pub: Zh. obshch. khimii, 1956, 26, No. 8, 2201 -
2209.

Abstract: The reactions of free radicals (forming in the
result of dissociation of alkylpenyltriazenes
and of dinitryl of aziisobutyric acid) with
natural rubber, synthetic polyisoprene and di-
vinyl polymer were studied in a wide range of
concentrations. The reactions of these polymers

High Polymer Inst. AN SSSR, Moscow.

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USSR/Chemistry of High Molecular Substances.

F

Abs Jour: Ref Zhur - Khimiya, No. 8, 1957, 27064.

with S_2Cl_2 were studied also. The polymer destruction takes place in all cases, the decomposition of the polymer chains occurs without any order, and the probability of a rupture at any monomer link of the polymer is the same. It is established that intramolecular vulcanization does not take place at the action of free radicals on polymers under the conditions, under which the reactions have been carried out. In the opinion of the authors, the destruction proceeds in two stages: 1/ tearing an H-atom away from the polymer with the formation of the polymer radical and 2/ dissociation of the polymer radical with the formation of the diene group-ation on the end of the chain and of the allyl radical. The authors arrive at the conclusion

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Sorption and chromatography of Ammonium O.V.
Samozetil S. V. Bremer and N. A. M. Gulyas, C. J.
(U.S.S.R.) 40, 401-4 (1966) (English translation). See
C 4 81 169

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